SYLLABUS

for

BACHELOR OF PHARMACY (B.Pharm) DEGREE PROGRAM

[Credit Based Semester System]

(Effective from the Academic year 2024–2025)

B. PHARM (SEMESTER - I)		
BP101T	BP101T HUMAN ANATOMY AND PHYSIOLOGY-I (Theory) 45	
functions o homeostati	s subject is designed to impart fundamental knowledge on the str f the various systems of the human body. It also helps in understa c mechanisms. The subject provides the basic knowledge re I the various disciplines of pharmacy.	nding both
	COURSE OUTCOMES	
CO 1	Understand the structure and function of the human body at the cellular, tissue, and organ levels.	molecular,
CO 2	Explain the physiological processes of different organ systems.	
CO 3	Describe the organization and function of the skeletal, cardiovascular, and nervous systems.	muscular,
CO 4	Analyze the mechanisms that maintain homeostasis in the body.	
CO 5	Apply anatomical and physiological knowledge in clinical context.	
COURSE CONTENT		
Unit I	Introduction to Human Body: Definition and scope of anatomy and physiology, levels of structural organization and body systems, basic life processes, homeostasis, basic anatomical terminology.	hours
	Cellular level of Organization: Structure and functions of cell, transport across cell membrane, cell division, cell junctions. General principles of cell communication, intracellular signaling pathway activation by extracellular signal molecule, Forms of intracellular signaling: a) Contact-dependent b) Paracrine c) Synaptic d) Endocrine Tissue level of ORGANIZATION: Classification of tissues,	
	structure, location and functions of epithelial, muscular and nervous and connective tissues.	
Unit II	Integumentary system: Structure and functions of skin Skeletal system: Divisions of skeletal system, types of bone, salient features and functions of bones of axial and appendicular skeletal system, Organization of skeletal muscle, physiology of muscle contraction, neuromuscular junction	nours

	Joints: Structural and functional classification, types of joints		
	movements and its articulation		
Unit III	Body fluids and blood: Body fluids, composition and functions of	10	
	blood, hemopoeisis, formation of hemoglobin, anemia,	hours	
	mechanisms of coagulation, blood grouping, Rh factors,	110415	
	transfusion, its significance and disorders of blood, Reticulo		
	endothelial system.		
	Lymphatic system: Lymphatic organs and tissues, lymphatic		
	vessels, lymph circulation and functions of lymphatic system.		
Unit IV	Peripheral nervous system: Classification of peripheral nervous	08	
	system: Structure and functions of sympathetic and	hours	
	parasympathetic nervous system. Origin and functions of spinal		
	and cranial nerves.		
	Special senses: Structure and functions of eye, ear, nose and		
	tongue and their disorders.		
Unit V	Cardiovascular system: Heart – anatomy of heart, blood	07	
	circulation, blood vessels, structure and functions of artery, vein	hours	
	and capillaries, elements of conduction system of heart and		
	heartbeat, its regulation by autonomic nervous system, cardiac		
	output, cardiac cycle. Regulation of blood pressure, pulse,		
	electrocardiogram and disorders of heart.		

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers' medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, River view, MI
- 4. Text book of Medical Physiology- Arthur C, Guyton and John.E. Hall. Miamisburg, OH, U.S.A.
- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A.
- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brother's medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brother's medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.
- 9. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 10. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 11. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterji

BP107P	HUMAN ANATOMY AND PHYSIOLOGY (Practical) 4 Hours/Week	
Scope: Pr	actical physiology is complimentary to the theoretical discussions in physiology.	
	allow the verification of physiological processes discussed in theory classes	
_	experiments on living tissue, intact animals or normal human beings. This is	
	r developing an insight on the subject.	
1	Study of compound microscope	
2	Microscopic study of epithelial and connective tissue	
3	Microscopic study of muscular and nervous tissue	
4	Identification of axial bones	
5	Identification of appendicular bones	
6	Introduction to hemocytometry	
7	Enumeration of white blood cell (WBC) count	
8	Enumeration of total red blood corpuscles (RBC) count	
9	Determination of bleeding time	
10	Determination of clotting time	
11	Estimation of haemoglobin content	
12	Determination of blood group	
13	Determination of erythrocyte sedimentation rate (ESR)	
14	Determination of heart rate and pulse rate	
15	Recording of blood pressure	

BP102T	PHARMACEUTICAL ANALYSIS (Theory)	45 Hours	
_	Scope: This course deals with the fundamentals of analytical chemistry and principles of electrochemical analysis of drugs		
	, ,		
	COURSE OUTCOMES		
CO 1	Minimize errors and enhance the accuracy and precision of	analytical	
	procedures.		
CO 2	Understand the principles of volumetric and electrochemical analy		
CO 3	Explain the various types of titrations used in pharmaceutical anal		
CO 4	Apply electrochemical techniques such as conductometry, potenti-	ometry and	
	polarography.		
CO 5	Apply analytical techniques to determine the quality and	purity of	
	pharmaceutical substances.		
	COURSE CONTENT		
Unit I	Pharmaceutical analysis- Definition and scope	10	
	a) Different techniques of analysis	hours	
	b) Methods of expressing concentration		
	c) Primary and secondary standards.		
	d) Preparation and standardization of various molar and normal		
	solutions- Oxalic acid, sodium hydroxide, hydrochloric acid,		
	sodium thiosulphate, sulphuric acid, potassium permanganate		
and ceric ammonium sulphate			
	Errors: Sources of errors, types of errors, methods of minimizing		
	errors, accuracy, precision and significant figures.		
	Pharmacopoeia, Sources of impurities in medicinal agents, limit		
TT. '- **	tests.	4.0	
Unit II	Acid base titration: Theories of acid base indicators, classification		
	of acid base titrations and theory involved in titrations of strong,	hours	
	weak, and very weak acids and bases, neutralization curves Non aqueous titration: Solvents, acidimetry and alkalimetry		
	titration and estimation of Sodium benzoate and Ephedrine HCl		
Unit III	Precipitation titrations: Mohr's method, Volhard's, Modified	10	
Onit iii	Volhard's, Fajans method, estimation of sodium chloride.	hours	
	i volitara o, rajano incatoa, coalitadon di soatam cinoriaci	nours	
	Complexometric titration: Classification, metal ion indicators.		
	Complexometric titration: Classification, metal ion indicators, masking and demasking reagents, estimation of Magnesium		

	Gravimetry: Principle and steps involved in gravimetric analysis.		
	Purity of the precipitate: co-precipitation and post precipitation,		
	Estimation of barium sulphate.		
	Basic Principles, methods and application of diazotization titration.		
Unit IV	Redox titrations	08	
	a) Concepts of oxidation and reduction	hours	
	b) Types of redox titrations (Principles and applications)		
	Cerimetry, Iodimetry, Iodometry, Bromatometry, Dichrometry,		
	Titration with potassium iodate		
Unit V	Electrochemical methods of analysis		
	Conductometry- Introduction, Conductivity cell, Conductometric	hours	
	titrations, applications.		
	Potentiometry - Electrochemical cell, construction and working of		
	reference (Standard hydrogen, silver chloride electrode and		
	calomel electrode) and indicator electrodes (metal electrodes and		
	glass electrode), methods to determine end point of potentiometric		
	titration and applications.		
	Polarography - Principle, Ilkovic equation, construction and		
	working of dropping mercury electrode and rotating platinum		
	electrode, applications		

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry
- 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 5. John H. Kennedy, Analytical chemistry principles
- 6. Indian Pharmacopoeia

PHARMACEUTICAL ANALYSIS (Practical) 4 Hours/Week **BP108P**

Scope: Practical physiology is complimentary to the theoretical discussions in physiology. Practical allow the verification of physiological processes discussed in theory classes through experiments on living tissue, intact animals or normal human beings. This is helpful for developing an insight on the subject.

Section	Topic
I	Limit Test of the following
1	Chloride
2	Sulphate
3	Iron
4	Arsenic
II	Preparation and standardization of
1	Sodium hydroxide
2	Sulphuric acid
3	Sodium thiosulfate
4	Potassium permanganate
5	Ceric ammonium sulphate
III	Assay of the following compounds along with Standardization of Titrant
1	Ammonium chloride by acid base titration
2	Ferrous sulphate by Cerimetry
3	Copper sulphate by Iodometry
4	Calcium gluconate by Complexometry
5	Hydrogen peroxide by Permanganometry
6	Sodium benzoate by non-aqueous titration
7	Sodium Chloride by precipitation titration
IV	Determination of Normality by electro-analytical methods
1	Conductometric titration of strong acid against strong base
2	Conductometric titration of strong acid and weak acid against strong base
3	Potentiometric titration of strong acid against strong base

BP103T	PHARMACEUTICS- I (Theory)	45 Hours		
_	Scope: This course is designed to impart a fundamental knowledge on the preparatory pharmacy			
with arts an	d science of preparing the different conventional dosage forms.			
	COURSE OUTCOMES			
CO 1	Understand the basic concepts of pharmaceutics and dosage form	ns.		
CO 2	Explain the principles of various pharmaceutical formulations.			
CO 3	Analyze the factors affecting drug absorption, distribution, meta excretion.	abolism, and		
CO 4	Understand the manufacturing processes of different dosage for	ns.		
CO 5	Apply knowledge of pharmaceutics to design and evaluate phadosage forms.	ırmaceutical		
	COURSE CONTENT			
Unit I	Historical background and development of profession o	of 10		
	pharmacy: History of profession of Pharmacy in India in relation	n hours		
	to pharmacy education, industry and organization, Pharmacy as	a		
	career, Pharmacopoeias: Introduction to IP, BP, USP and Extra			
	Pharmacopoeia.			
	Dosage forms: Introduction to dosage forms, classification and			
	definitions			
	Prescription: Definition, Parts of prescription, handling of Prescription and Errors in prescription.			
	Posology: Definition, Factors affecting posology. Pediatric dose			
TI!4 TT	calculations based on age, body weight and body surface area.	1 10		
Unit II	Pharmaceutical calculations: Weights and measures – Imperia			
	& Metric system, Calculations involving percentage solutions			
	alligation, proof spirit and isotonic solutions based on freezing point and molecular weight.	8		
	Powders: Definition, classification, advantages and	d		
	disadvantages, Simple & compound powders – officia			
	preparations, dusting powders, effervescent, efflorescent and			
	hygroscopic powders, eutectic mixtures. Geometric dilutions.			
	Liquid dosage forms: Advantages and disadvantages of liquid	d		
	dosage forms. Excipients used in formulation of liquid dosag			
	forms. Solubility enhancement techniques			

Unit III	Monophasic liquids: Definitions and preparations of Gargles,	10		
	Mouthwashes, Throat Paint, Eardrops, Nasal drops, Enemas,	hours		
	Syrups, Elixirs, Liniments and Lotions.			
	Biphasic liquids:			
	Suspensions: Definition, advantages and disadvantages,			
	classifications, Preparation of suspensions; Flocculated and			
	Deflocculated suspension & stability problems and methods to			
	overcome.			
	Emulsions: Definition, classification, emulsifying agent, test for			
	the identification of type of Emulsion, Methods of preparation &			
	stability problems and methods to overcome.			
Unit IV	Suppositories: Definition, types, advantages and disadvantages, 08			
	types of bases, methods of preparations. Displacement value & its	hours		
	calculations, evaluation of suppositories.			
	Pharmaceutical incompatibilities: Definition, classification,			
	physical, chemical and therapeutic incompatibilities with			
	examples.			
Unit V	Semisolid dosage forms: Definitions, classification, mechanisms	07		
	and factors influencing dermal penetration of drugs. Preparation	hours		
	of ointments, pastes, creams and gels. Excipients used in semi solid			
	dosage forms. Evaluation of semi solid dosages forms			

- 1. H.C. Ansel et al., pharmaceutical dosage form and drug delivery system, Lippincott Williams and Walkins, New Delhi.
- 2. Carter S.J., Cooper and Gunn's-Dispensing for pharmaceutical students, CBS publishers, New Delhi.
- 3. M.E. Aulton, Pharmaceutics, The science & dosage form design, Churchill Livingstone, Edinburgh.
- 4. Indian pharmacopoeia.
- 5. Lachmann, Theory and practice of industrial pharmacy, Lea & Febiger Publisher, The University of Michigan.
- 6. Alfonso R. Gennaro, Remington: The science and practice of pharmacy, Lippincott Williams, New Delhi.
- 7. Carter S.J., Cooper and Gunn's, Tutorial pharmacy, CBS Publications, New Delhi.
- 8. E.A. Rawlins, Bentley's text book of pharmaceutics, English Language Book Society, Elsevier Health Sciences, USA.
- 9. Isaac Ghebre Sellassie, Pharmaceutical pelletization technology, Marcel Dekker, INC, New York.
- 10. Dilip M. Parikh, Handbook of pharmaceutical granulation technology, Marcel Dekker, INC, New
- 11. Francoise Nieloud and Gilberte Marti-Mestres, Pharmaceutical emulsions and suspensions, Marcel Dekker, INC, New York.

BP109P	PHARMACEUTI	CS- I (Practical)	4 Hours/Week
S.No.	Category	a) Syrup IP'66	
		b) Compound syrup of Ferro	us Phosphate BPC'68
1	Syrups	a) Piperazine citrate elixir	
		b) Paracetamol pediatric elix	xir
2	Elixirs	a) Terpin Hydrate Linctus IP	2'66
		b) Iodine Throat Paint (Man	dles Paint)
3	Linctus	a) Strong solution of ammor	ium acetate
		b) Cresol with soap solution	
		c) Lugol's solution	
4	Solutions	a) Calamine lotion	
		b) Magnesium Hydroxide m	ixture
		c) Aluminium Hydroxide gel	
5	Suspensions	a) Turpentine Liniment	
		b) Liquid paraffin emulsion	
6	Emulsions	a) ORS powder (WHO)	
		b) Effervescent granules	
		c) Dusting powder	
		d) Divided powders	
7	Powders and Granules	a) Glycero gelatin supposito	ry
		b) Cocoa butter suppository	
		c) Zinc Oxide suppository	
8	Suppositories	a) Sulphur ointment	
		b) Non-staining-iodine oin	itment with methyl
		salicylate	
		c) Carbopol gel	
9	1) 11 11 11 81		
		b) Chlorhexidine Mouthwas	h
10	Gargles and mouthwashes	a) Syrup IP'66	
		b) Compound syrup of Ferro	us Phosphate BPC'68

DD 40 :-			
BP104T	PHARMACEUTICAL INORGANIC CHEMISTRY (Theory)	45 Hours	
Scope: This subject deals with the monographs of inorganic drugs and pharmaceuticals.			
	COURSE OUTCOMES		
60.4			
CO 1	Understand the classification, nomenclature, and properties of compounds.	inorganic	
CO 2	Explain the preparation, uses, and analysis of various inorganic pharcompounds.	rmaceutical	
CO 3	Analyse the importance of inorganic compounds in pharmacy and medi	cine.	
CO 4	Understand the concepts of qualitative and quantitative analysis of compounds.	f inorganic	
CO 5	Understand the principles and applications of radiopharmaceuticals	in medical	
	practice.		
	COURSE CONTENT		
Unit I	Impunities in pharmacoutical substances History of		
Uniti	Impurities in pharmaceutical substances: History of Pharmacopoeia, Sources and types of impurities, principle	10	
	involved in the limit test for Chloride, Sulphate, Iron, Arsenic, Lead	hours	
	and Heavy metals, modified limit test for Chloride and Sulphate		
	General methods of preparation, assay for the compounds		
	superscripted with asterisk (*), properties and medicinal uses of		
	inorganic compounds belonging to the following classes		
Unit II	Acids, Bases and Buffers: Buffer equations and buffer capacity in	10	
	general, buffers in pharmaceutical systems, preparation, stability,	hours	
	buffered isotonic solutions, measurements of tonicity, calculations		
	and methods of adjusting isotonicity.		
	Major extra and intracellular electrolytes: Functions of major		
	physiological ions, Electrolytes used in the replacement therapy:		
	Sodium chloride*, Potassium chloride, Calcium gluconate* and		
	Oral Rehydration Salt (ORS), Physiological acid base balance.		
	Dental products: Dentifrices, role of fluoride in the treatment of		
	dental caries, Desensitizing agents, Calcium carbonate, Sodium fluoride, and Zinc eugenol cement.		
Unit III	Gastrointestinal agents	10	
	Acidifiers: Ammonium chloride* and Dil. HCl	hours	
	Antacid: Ideal properties of antacids, combinations of		
	antacids, Sodium Bicarbonate*, Aluminum hydroxide gel,		
	Magnesium hydroxide mixture		

	Cathartics: Magnesium sulphate, Sodium orthophosphate, Kaolin			
	and Bentonite			
	Antimicrobials: Mechanism, classification, Potassium			
	permanganate, Boric acid, Hydrogen peroxide*, Chlorinated lime*,			
	Iodine and its preparations			
Unit IV	Miscellaneous compounds	08		
	Expectorants: Potassium iodide, Ammonium chloride*. Emetics:	hours		
	Copper sulphate*, Sodium potassium tartarate Haematinics:			
	Ferrous sulphate*, Ferrous gluconate			
	Poison and Antidote: Sodium thiosulphate*, Activated charcoal,			
	Sodium nitrite333			
	Astringents: Zinc Sulphate, Potash Alum			
Unit V	Radiopharmaceuticals: Radio activity, Measurement of	07		
	radioactivity, Properties of α , β , γ radiations, Half-life, radio	hours		
	isotopes and study of radio isotopes - Sodium iodide I ¹³¹ , Storage			
	conditions, precautions & pharmaceutical application of			
	radioactive substances.			

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London, 4th edition.
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry, 3rd Edition
- 4. M.L Schroff, Inorganic Pharmaceutical Chemistry
- 5. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 6. Anand & Chatwal, Inorganic Pharmaceutical Chemistry
- 7. Indian Pharmacopoeia

BP110P	PHARMACEUTICAL INORGANIC CHEMISTRY (Practical) 4 Hours/Week		
S.No.	Category	Experiment	
1	Limit tests for following	a) Limit test for Chlorides and Sulphates	
	ions	b) Modified limit test for Chlorides and Sulphates	
		c) Limit test for Iron	
		d) Limit test for Heavy metals	
		e) Limit test for Lead	
		f) Limit test for Arsenic	
2	Identification test	g) Magnesium hydroxide	
		h) Ferrous sulphate	
		i) Sodium bicarbonate	
		j) Calcium gluconate	
		k) Copper sulphate	
3	Test for purity	l) Swelling power of Bentonite	
		m) Neutralizing capacity of aluminium hydroxide gel	
		n) Determination of potassium iodate and iodine in	
		potassium iodide	
4	Preparation of inorganic	o) Boric acid	
	pharmaceuticals	p) Potash alum	
		q) Ferrous sulphate	

BP105T	COMMUNICATION SKILLS (Theory)	30 Hours
Scope: This course will prepare the young pharmacy student to interact effectively with doct nurses, dentists, physiotherapists and other health workers. At the end of this course the stud will get the soft skills set to work cohesively with the team as a team player and will add value the pharmaceutical business.		
	COURSE OUTCOMES	
CO 1	Understand the fundamentals of effective communication.	
CO 2	Explain the importance of communication in professional an contexts.	d personal
CO 3	Develop skills in verbal and non-verbal communication.	
CO 4	Analyze various barriers to communication and strategies to over	
CO 5	Apply communication skills in pharmacy practice, includi	ng patient
	counselling and professional interactions.	
	COURSE CONTENT	
Unit I	Communication Skills: Introduction, Definition, The Importance of Communication, The Communication Process – Source Message, Encoding, Channel, Decoding, Receiver, Feedback Context Barriers to communication: Physiological Barriers, Physica Barriers, Cultural Barriers, Language Barriers, Gender Barriers Interpersonal Barriers, Psychological Barriers, Emotional barriers Perspectives in Communication: Introduction, Visual Perception Language, Other factors affecting our perspective - Passe Experiences, Prejudices, Feelings, Environment	l s,
Unit II	Elements of Communication: Introduction, Face to Face Communication - Tone of Voice, Body Language (Non-verbal communication), Verbal Communication, Physical Communication Communication Styles: Introduction, The Communication Styles Matrix with example for each -Direct Communication Style Spirited Communication Style, Systematic Communication Style Considerate Communication Style	
Unit III	Basic Listening Skills: Introduction, Self-Awareness, Active Listening, Becoming an Active Listener, Listening in Difficult Situations	

	Effective Written Communication: Introduction, When and		
	When Not to Use Written Communication - Complexity of the		
	Topic, Amount of Discussion' Required, Shades of Meaning, Formal		
	Communication		
	Writing Effectively: Subject Lines, Put the Main Point First, Know		
	Your Audience, Organization of the Message		
Unit IV	Interview Skills: Purpose of an interview, Do's and Dont's of an 5 hours		
	interview		
	Giving Presentations: Dealing with Fears, planning your		
	Presentation, Structuring Your Presentation, Delivering Your		
	Presentation, Techniques of Delivery		
Unit V	Group Discussion: Introduction, Communication skills in group	4 hours	
	discussion, Do's and Dont's of group discussion		

- 1. Basic communication skills for Technology, Andreja. J. Ruther Ford, 2nd Edition, Pearson Education, 2011
- 2. Communication skills, Sanjay Kumar, Pushpalata, 1stEdition, Oxford Press, 2011
- 3. Organizational Behaviour, Stephen .P. Robbins, 1stEdition, Pearson, 2013
- 4. Brilliant- Communication skills, Gill Hasson, 1stEdition, Pearson Life, 2011
- 5. The Ace of Soft Skills: Attitude, Communication and Etiquette for success, Gopala Swamy Ramesh, 5thEdition, Pearson, 2013
- 6. Developing your influencing skills, Deborah Dalley, Lois Burton, Margaret, Green hall, 1st Edition Universe of Learning LTD, 2010
- 7. Communication skills for professionals, Konar nira, 2ndEdition, New arrivals –
- 8. PHI, 2011
- 9. Personality development and soft skills, Barun K Mitra, 1stEdition, Oxford Press, 2011
- 10. Soft skill for everyone, Butter Field, 1st Edition, Cengage Learning india pyt.ltd,
- 11. Soft skills and professional communication, Francis Peters SJ, 1stEdition, Mc Graw Hill Education, 2011
- 12. Effective communication, John Adair, 4thEdition, Pan Mac Millan, 2009
- 13. Bringing out the best in people, Aubrey Daniels, 2ndEdition, Mc Graw Hill, 1999

BP111P	COMMUNICAT	2 Hours/Week	
S.No.	Category	Experiment	
1	Basic Communication	- Meeting People	
		- Asking Questions	
		- Making Friends	
		- What did you do?	
		- Do's and Don'ts	
2	Pronunciations	- Pronunciation (Consonant Sounds)	
		- Pronunciation and Nouns	
		- Pronunciation (Vowel Sounds)	
3	Advanced Learning	- Listening Comprehension /	Direct and Indirect
		Speech	
		- Figures of Speech	
		- Effective Communication	
		- Writing Skills	
		- Effective Writing	
		- Interview Handling Skills	
		- E-Mail etiquette	
		- Presentation Skills	

BP106RBT	REMEDIAL BIOLOGY (Theory)	30 Hours
-	rn and understand the components of living world, structure and funct	ional system
of plant and a	nimal kingdom.	
	COURSE OUTCOMES	
CO 1	Understand the classification and key characteristics of the five kinge	doms of life.
CO 2	Comprehend the basic components and functions of plant a	and animal
	anatomy and physiology with a focus on human systems.	
CO 3	Gain knowledge of various human body systems, including	circulatory,
	digestive, respiratory, excretory, nervous, endocrine and reproductive	
CO 4		rowth and
	development.	
CO 5	Grasp the structure and function of cells and their organelles, as well	as the types
	and functions of tissues and the process of cell division.	
	COURSE CONTENT	
Unit I	Living world:	7 hours
	Definition and characters of living organisms, Diversity in the	
	living world, Binomial nomenclature, Five kingdoms of life and	
	basis of classification. Salient features of Monera, Potista, Fungi	,
	Animalia and Plantae, Virus,	
	Morphology of Flowering plants Morphology of different parts of flowering plants – Root, stem	
	inflorescence, flower, leaf, fruit, seed.	,
	General Anatomy of Root, stem, leaf of monocotyledons &	,
	Dicotylidones.	
Unit II	Body fluids and circulation	7 h c
	Composition of blood, blood groups, coagulation of blood	7 hours
	Composition and functions of lymph; Human circulatory system	
	Structure of human heart and blood vessels; Cardiac cycle, cardiac	
	output and ECG	
	Digestion and Absorption	
	Human alimentary canal and digestive glands; Role of digestive	
	enzymes; Digestion, absorption and assimilation of digested food	;
	Breathing and respiration; Human respiratory system	
	Mechanism of breathing and its regulation; Exchange of gases	
	transport of gases and regulation of respiration; Respiratory	7
	volumes	

Unit III	Excretory products and their elimination	7 hours
	Modes of excretion; Human excretory system- structure and	
	function; Urine formation; Rennin angiotensin system	
	Neural control and coordination	
	Definition and classification of nervous system; Structure of a	
	neuron; Generation and conduction of nerve impulse;	
	Structure of brain and spinal cord; Functions of cerebrum,	
	cerebellum, hypothalamus and medulla oblongata;	
	Chemical coordination and regulation	
	Endocrine glands and their secretions; Functions of hormones	
	secreted by endocrine glands	
	Human reproduction	
	Parts of female reproductive system; Parts of male	
	reproductive system; Spermatogenesis and Oogenesis; Menstrual	
	cycle	
Unit IV	Plants and mineral nutrition:	5 hours
	Essential mineral, macro and micronutrients; Nitrogen	
	metabolism, Nitrogen cycle, biological nitrogen fixation	
	Photosynthesis	
	Autotrophic nutrition, photosynthesis, Photosynthetic pigments,	
	Factors affecting photosynthesis.	
Unit V	Plant respiration: Respiration, glycolysis, fermentation	4 hours
	(anaerobic).	
	Plant growth and development	
	Phases and rate of plant growth, Condition of growth,	
	Introduction to plant growth regulators Cell - The unit of life	
	Structure and functions of cell and cell organelles. Cell division	
	Tissues	
	Definition, types of tissues, location and functions.	

Text Books

- 1. Text book of Biology by S. B. Gokhale
- 2. A Text book of Biology by Dr. Thulajappa and Dr. Seetaram.

Reference Books

- 3. A Text book of Biology by B.V. Sreenivasa Naidu
- 4. A Text book of Biology by Naidu and Murthy
- 5. Botany for Degree students By A.C. Dutta.
- 6. Outlines of Zoology by M. Ekambaranatha ayyer and T. N. Ananthakrishnan.
- 7. A manual for pharmaceutical biology practical by S.B. Gokhale and C. K. Kokate

BP112RBP	REMEDIAL BIOLOGY (Practical)		30 Hours	
S.No.	Category	Expe	eriment	
1	Introduction to	a)	Study of Microscope	
	experiments in biology	b)	Section cutting technique	ies
		c)	Mounting and staining	
		d)	Permanent slide prepar	ation
2	Study of cell and its inclusion	ns		
3	Study of stem, root, leaf, seed, fruit, flower, and their modifications			
4	Detailed study of frog by using computer models			
5	Microscopic study and identification of tissues pertinent to stem, root, leaf,			
	seed, fruit, and flower			
6	Identification of bones			
7	Determination of blood group			
8	Determination of blood pressure			
9	Determination of tidal volume			

- 1. Practical human anatomy and physiology. by S.R. Kale and R.R. Kale.
- 2. A Manual of pharmaceutical biology practical by S.B. Gokhale, C.K. Kokate and S.P. Shriwastava.
- 3. Biology practical manual according to National core curriculum. Biology forum of Karnataka. Prof.M.J.H. Shafi

BP106RMT	REMEDIAL MATHEMATICS (Theory)	30 Hours
Scope: This is	an introductory course in mathematics. This subject deals with the in	troduction to
Partial fraction	n, Logarithm, matrices and Determinant, Analytical geometry, Calculus	s, differential
equation and l	Laplace transform.	
	COURSE OUTCOMES	
CO 1	Understand the fundamental concepts of mathematics relevant to	pharmacy.
CO 2	Explain the applications of calculus, algebra, and trigo pharmaceutical sciences.	nometry in
CO 3	Analyze mathematical models used in pharmacoking	netics and
	pharmacodynamics.	
CO 4	Solve mathematical problems related to drug formulation and dos	sing.
CO 5	Apply mathematical principles in pharmaceutical research and practical research and practic	actice.
	COURSE CONTENT	
Unit I	Partial fraction	6 hours
	Introduction, Polynomial, Rational fractions, Proper and Improper fractions, Partial fraction, Resolving into Partial fraction, Application of Partial Fraction in Chemical Kinetics and Pharmacokinetics Logarithms Introduction, Definition, Theorems/Properties of logarithms, Common logarithms, Characteristic and Mantissa, worked	o nom s
	examples, application of logarithm to solve pharmaceutical problems. Function:	
	Real Valued function, Classification of real valued functions, Limits and continuity: Introduction, Limit of a function, Definition of limit of a function ($\in -\delta$ definition), $\lim_{x \to a} \frac{x^n - a^n}{x - a} = na^{n-1}$, $\lim_{\theta \to 0} \frac{\sin \theta}{\theta} = 1$,	
Unit II	Matrices and Determinant:	6 hours
	Introduction matrices, Types of matrices, Operation on matrices, Transpose of a matrix, Matrix Multiplication, Determinants, Properties of determinants, Product of determinants, Minors and co-Factors, Adjoint or adjugate of a square matrix, Singular and nonsingular matrices, Inverse of a matrix, Solution of system of linear of equations using matrix method, Cramer's rule, Characteristic equation and roots of a square matrix, Cayley–Hamilton theorem, Application of Matrices in solving Pharmacokinetic equations.	

Unit III	Calculus	6 hours	
	Differentiation: Introductions, Derivative of a function, Derivative		
	of a constant, Derivative of a product of a constant and a function,		
	Derivative of the sum or difference of two functions, Derivative of		
	the product of two functions (product formula), Derivative of the		
	quotient of two functions (Quotient formula) - Without Proof,		
	Derivative of x n w.r.tx, where n is any rational number, Derivative		
	of ex,, Derivative of loge x , Derivative of a x ,Derivative of		
	trigonometric functions from first principles (without Proof),		
	Successive Differentiation, Conditions for a function to be a		
	maximum or a minimum at a point. Application.		
Unit IV	Analytical Geometry	6 hours	
	Introduction: Signs of the Coordinates, Distance formula,		
	Straight Line: Slope or gradient of a straight line, Conditions for		
	parallelism and perpendicularity of two lines, Slope of a line joining		
	two points, Slope – intercept form of a straight line		
	Integration:		
	Introduction, Definition, Standard formulae, Rules of integration,		
	Method of substitution, Method of Partial fractions, Integration by		
	parts, definite integrals, application.		
Unit V	Differential Equations: Some basic definitions, Order and degree,	6 hours	
	Equations in separable form, Homogeneous equations, Linear		
	Differential equations, Exact equations,		
	Application in solving Pharmacokinetic equations		
	Laplace Transform: Introduction, Definition, Properties of		
	Laplace transform, Laplace Transforms of elementary functions,		
	Inverse Laplace transforms, Laplace transform of derivatives,		
	Application to solve Linear differential equations,		
	Application in solving Chemical kinetics and		
	Pharmacokinetics equations		

- 1. Differential Calculus by Shanthinarayan
- 2. Pharmaceutical Mathematics with application to Pharmacy by Panchaksharappa Gowda D.H.
- 3. Integral Calculus by Shanthinarayan
- 4. Higher Engineering Mathematics by Dr.B.S.Grewal

B. PHARM (SEMESTER - II)				
BP201T	HUMAN ANATOMY AND PHYSIOLOGY-II (Theory)			
functions o	s subject is designed to impart fundamental knowledge on the str of the various systems of the human body. It also helps in understa of mechanisms. The subject provides the basic knowledge re the various disciplines of pharmacy.	nding both		
	COURSE OUTCOMES			
CO 1	Understand the anatomy and physiology of various body systems the nervous, endocrine, cardiovascular, respiratory, digestive, unreproductive systems.	_		
CO 2	Explain the biochemical and physiological processes underlying the of different organs and systems.	he function		
CO 3	Describe the mechanisms of homeostasis and how they maintain body equilibrium. Explain the introduction to genetics and discuss genetic pattern of inheritance			
CO 4	Analyze the physiological basis of health and disease states.			
CO 5	Apply knowledge of human anatomy and physiology in clinical contexts.			
	COURSE CONTENT			
Unit I	Nervous system Organization of nervous system, neuron, neuroglia, classification and properties of nerve fibre, electrophysiology, action potential, nerve impulse, receptors, synapse, neurotransmitters. Central nervous system: Meninges, ventricles of brain and cerebrospinal fluid. Structure and functions of brain (cerebrum, brain stem, cerebellum), spinal cord (gross structure, functions of afferent and efferent nerve tracts, reflex activity)	10 hours		
Unit II	Anatomy of GI Tract with special reference to anatomy and functions of stomach, (Acid production in the stomach, regulation of acid production through parasympathetic nervous system, pepsin role in protein digestion) small intestine and large intestine, anatomy and functions of salivary glands, pancreas and liver, movements of GIT, digestion and absorption of nutrients and disorders of GIT. Energetics Formation and role of ATP, Creatinine Phosphate and BMR.	06 hours		

Unit III	Respiratory system Anatomy of respiratory system with special reference to anatomy of lungs, mechanism of respiration, regulation of respiration Lung Volumes and capacities transport of respiratory gases, artificial respiration, and resuscitation methods. Urinary system Anatomy of urinary tract with special reference to anatomy of kidney and nephrons, functions of kidney and urinary tract, physiology of urine formation, micturition reflex and role of kidneys in acid base balance, role of RAS in kidney and disorders of kidney.	10 hours
Unit IV	Endocrine system Classification of hormones, mechanism of hormone action, structure and functions of pituitary gland, thyroid gland, parathyroid gland, adrenal gland, pancreas, pineal gland, thymus and their disorders.	10 hours
Unit V	Reproductive system Anatomy of male and female reproductive system, Functions of male and female reproductive system, sex hormones, physiology of menstruation, fertilization, spermatogenesis, oogenesis, pregnancy and parturition Introduction to genetics Chromosomes, genes and DNA, protein synthesis, genetic pattern of inheritance	09 hours

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co,Riverview,MI USA
- 4. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A.
- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brothers medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brothers medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.
- 9. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 10. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 11. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterrje, Academic Publishers Kolkata

BP207P	HUMAN ANATOMY AND PHYSIOLOGY (Practical)	4 Hours/Week

Scope: Practical physiology is complimentary to the theoretical discussions in physiology. Practicals allow the verification of physiological processes discussed in theory classes through experiments on living tissue, intact animals or normal human beings. This is helpful for developing an insight on the subject.

S.No.	Topic
1	To study the integumentary and special senses using specimens, models, etc.
2	To study the nervous system using specimens, models, etc.
3	To study the endocrine system using specimens, models, etc.
4	To demonstrate the general neurological examination
5	To demonstrate the function of the olfactory nerve
6	To examine the different types of taste
7	To demonstrate visual acuity
8	To demonstrate reflex activity
9	Recording of body temperature
10	To demonstrate positive and negative feedback mechanisms
11	Determination of tidal volume and vital capacity
12	Study of digestive, respiratory, cardiovascular, urinary, and reproductive
	systems with models, charts, and specimens
13	Recording of basal metabolic index
14	Study of family planning devices and pregnancy diagnosis tests
15	Demonstration of total blood count by cell analyzer
16	Permanent slides of vital organs and gonads

BP202T PHARMACEUTICAL ORGANIC CHEMISTRY -I (Theory) 45 Hours
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Scope: This subject deals with classification and nomenclature of simple organic compounds, structural isomerism, intermediates forming in reactions, important physical properties, reactions and methods of preparation of these compounds. The syllabus also emphasizes on mechanisms and orientation of reactions.

	COURSE OUTCOMES
CO 1	Gain foundational knowledge of organic chemistry concepts including classification, nomenclature, and isomerism.
	Explain the mechanisms of organic reactions, including electrophilic and
CO 2	nucleophilic substitution, elimination, and addition reactions.
CO 3	Analyse the various naming reactions and significance of qualitative tests in
	organic chemistry.
CO 4	Understand the synthesis and reactions of functional groups important in drug
CO 4	design and development.
CO 5	Apply organic chemistry principles to draw the structure and uses of organic
CO 5	compounds.

COURSE CONTENT

General methods of preparation and reactions of compounds superscripted with asterisk (*) to be explained.

To emphasize on definition, types, classification, principles/mechanisms, applications, examples and differences.

Unit I	Classification, nomenclature and isomerism	07 hours
	Classification of Organic Compounds	or nours
	Common and IUPAC systems of nomenclature of organic	
	compounds (up to 10 Carbons open chain and carbocyclic	
	compounds); Structural isomerisms in organic compounds	
Unit II	Alkanes*, Alkenes* and Conjugated dienes*	10 hours
	SP3 hybridization in alkanes, Halogenation of alkanes, uses of	10 nours
	paraffins. Stabilities of alkenes, SP2 hybridization in alkenes	
	E1 and E2 reactions - kinetics, order of reactivity of alkyl	
	halides, rearrangement of carbocations, Saytzeffs orientation	
	and evidences. E1 verses E2 reactions, Factors affecting E1 and	
	E2 reactions. Ozonolysis, electrophilic addition reactions of	
	alkenes, Markownikoff's orientation, free radical addition	
	reactions of alkenes, Anti Markownikoff's orientation. Stability	
	of conjugated dienes, Diel-Alder, electrophilic addition, free	
	radical addition reactions of conjugated dienes, allylic	
	rearrangement	

Unit III	Alkyl halides*	10 hours
	SN1 and SN2 reactions - kinetics, order of reactivity of alkyl	
	halides, stereochemistry and rearrangement of carbocations.	
	SN1 versus SN2 reactions, Factors affecting SN1 and SN2	
	reactions.	
	Structure and uses of ethylchloride, Chloroform,	
	trichloroethylene, tetrachloroethylene, dichloromethane,	
	tetrachloromethane and iodoform.	
	Alcohols* Qualitative tests, Structure and uses of Ethyl alcohol,	
	Methyl alcohol, chlorobutanol, Cetosteryl alcohol, Benzyl alcohol,	
	Glycerol, Propylene glycol	
Unit IV	Carbonyl compounds* (Aldehydes and ketones)	10 hours
	Nucleophilic addition, Electromeric effect, aldol condensation,	
	Crossed Aldol condensation, Cannizzaro reaction, Crossed	
	Cannizzaro reaction, Benzoin condensation, Perkin	
	condensation, qualitative tests, Structure and uses of	
	Formaldehyde, Paraldehyde, Acetone, Chloral hydrate,	
	Hexamine, Benzaldehyde, Vanilin, Cinnamaldehyde.	
Unit V	Carboxylic acids*	09
	Acidity of carboxylic acids, effect of substituents on acidity,	hours
	inductive effect and qualitative tests for carboxylic acids, amide	
	and ester Structure and Uses of Acetic acid, Lactic acid, Tartaric	
	acid, Citric acid, Succinic acid. Oxalic acid, Salicylic acid, Benzoic	
	acid, Benzyl benzoate, Dimethyl phthalate, Methyl salicylate and	
	Acetyl salicylic acid	
	Aliphatic amines*	
	Basicity, effect of substituent on Basicity. Qualitative test,	
	Structure and uses of Ethanolamine, Ethylenediamine,	
	Amphetamine.	

- Organic Chemistry by Morrison and Boyd 1.
- Organic Chemistry by I.L. Finar, Volume-I 2.
- Textbook of Organic Chemistry by B.S. Bahl & Arun Bahl. 3.
- 4. Organic Chemistry by P.L. Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K. Vishnoi.
- Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz. 8.
- Reaction and reaction mechanism by Ahluwaliah/Chatwal.

BP208P	PHARMACEUTICAL ORGANIC CHEMISTRY -I (Practical)	4 Hours/Week
Systemat	tic qualitative analysis of unknown organic compounds like	
1	Preliminary test: Color, odour, aliphatic/aromatic compounds,	saturation and
	unsaturation, etc.	
2	Detection of elements like Nitrogen, Sulphur & Halogen by Last	saigne's test
3	Solubility test	
4	Functional group test like Phenols, Amides/ Urea, Carbohydrat	es, Amines,
	Carboxylic acids, Aldehydes and Ketones, Alcohols, Esters, Aror	natic and
	Halogenated Hydrocarbons, Nitro compounds and Anilides.	
5	Melting point/Boiling point of organic compounds	
6	Identification of the unknown compound from the literature us	ing melting point/
	boiling point.	
7	Preparation of the derivatives and confirmation of the unknown	n compound by
	melting point/ boiling point.	
8	Minimum 5 unknown organic compounds to be analysed system	natically.
Preparation of suitable solid derivatives from organic compounds		
Construction of molecular models		

		I SEMESTER	
BP203T	BIOCHEMISTRY (Theory)	45 Hours	
Scope: Bio	chemistry deals with complete understanding of the molecul	ar levels of the	
chemical p	chemical process associated with living cells. The scope of the subject is providing		
biochemica	biochemical facts and the principles to understand metabolism of nutrient molecules in		
physiologic	cal and pathological conditions.		
	COURCE OUTCOMEC		
	COURSE OUTCOMES		
CO 1	Understand the structure and function of biomolecules, including	ng proteins,	
	carbohydrates, lipids, and nucleic acids.		
CO 2	Explain the metabolic pathways and their regulation in the hum	ıan body.	
CO 3	Understand the pathways and regulation of amino acid m	etabolism, lipid	
	metabolism and their roles in health and disease.		
CO 4	Understand the pathways and regulatory mechanisms of	of nucleic acid	
	metabolism, including DNA replication, transcription, and repair	r mechanisms.	
CO 5	Analyze the role of enzymes in catalyzing biochemical reactions	S	
	COURSE CONTENT		
	COURSE CONTENT		
Unit I	Biomolecules	08 hours	
	Introduction, classification, chemical nature and biological role		
	of carbohydrate, lipids, nucleic acids, amino acids and proteins	i.	
	Bioenergetics		
	Concept of free energy, endergonic and exergonic reaction		
	Relationship between free energy, enthalpy and entropy; Redo		
	potential. Energy rich compounds; classification; biologica	1	
	significances of ATP and cyclic AMP		
Unit II	Carbohydrate metabolism	10	
	Glycolysis – Pathway, energetics and significance Citric acid	**	
	cycle- Pathway, energetics and significance; HMP shunt and it	3	
	significance; Glucose-6-Phosphate dehydrogenase (G6PD		
	deficiency; Glycogen metabolism Pathways and glycoger storage diseases (GSD) Gluconeogenesis- Pathway and it		
	significance;	5	
	Hormonal regulation of blood glucose level and Diabete	c	
	mellitus	5	
	Biological oxidation		
	Electron transport chain (ETC) and its mechanism. Oxidative	e	
	phosphorylation & its mechanism and substrate		
	phosphorylation		
	Inhibitors ETC and oxidative phosphorylation/Uncouplers		

level

Unit III

Lipid metabolism

10 hours

	β-Oxidation of saturated fatty acid (Palmitic acid)	
	Formation and utilization of ketone bodies; ketoacidosis De novo	
	synthesis of fatty acids (Palmitic acid) Biological significance of	
	cholesterol and conversion of cholesterol into bile acids, steroid	
	hormone and vitamin D Disorders of lipid metabolism:	
	Hypercholesterolemia, atherosclerosis, fatty liver and obesity.	
	Amino acid metabolism	
	General reactions of amino acid metabolism: Transamination,	
	deamination & decarboxylation, urea cycle and its disorders	
	Catabolism of phenylalanine and tyrosine and their metabolic	
	disorders (Phenyketonuria, Albinism, alkeptonuria,	
	tyrosinemia) Synthesis and significance of biological substances;	
	5-HT, melatonin, dopamine, noradrenaline, adrenaline	
	Catabolism of heme; hyperbilirubinemia and jaundice	
Unit IV	Nucleic acid metabolism and genetic information transfer	10 hours
	Biosynthesis of purine and pyrimidine nucleotides;	
	Catabolism of purine nucleotides and Hyperuricemia and Gout	
	disease; Organization of mammalian genome; Structure of DNA	
	and RNA and their functions; DNA replication (semi conservative	
	model); Transcription or RNA synthesis; Genetic code,	
	Translation or Protein synthesis and inhibitors	
Unit V	Enzymes	09
	Introduction, properties, nomenclature and IUB classification of	hours
	enzymes; Enzyme kinetics (Michaelis plot, Line Weaver Burke	
	plot); Enzyme inhibitors with examples; Regulation of enzymes:	
1	1	
	enzyme induction and repression, allosteric enzymes regulation	
	enzyme induction and repression, allosteric enzymes regulation Therapeutic and diagnostic applications of enzymes and	

- 1. Principles of Biochemistry by Lehninger.
- 2. Harper's Biochemistry by Robert K. Murry, Daryl K. Granner and Victor W. Rodwell.
- 3. Biochemistry by Stryer.
- 4. Biochemistry by D. Satyanarayan and U. Chakrapani
- 5. Textbook of Biochemistry by Rama Rao.
- 6. Textbook of Biochemistry by Deb.
- 7. Outlines of Biochemistry by Conn and Stumpf
- 8. Practical Biochemistry by R.C. Gupta and S. Bhargavan.
- 9. Introduction of Practical Biochemistry by David T. Plummer. (3rd Edition)
- 10. Practical Biochemistry for Medical students by Rajagopal and Ramakrishna.
- 11. Practical Biochemistry by Harold Varley.

BP209P	PHARMACEUTICAL BIOCHEMISTRY (Practical)	4 Hours/Week
S.No.	Experiment	
1	Qualitative analysis of carbohydrates (Glucose, Fructose	e, Lactose, Maltose,
	Sucrose, and starch)	
2	Identification tests for Proteins (albumin and Casein)	
3	Quantitative analysis of reducing sugars (DNSA method) and Proteins
	(Biuret method)	
4	Qualitative analysis of urine for abnormal constituents	
5	Determination of blood creatinine	
6	Determination of blood sugar	
7	Determination of serum total cholesterol	
8	Preparation of buffer solution and measurement of pH	
9	Study of enzymatic hydrolysis of starch	
10	Determination of Salivary amylase activity	
11	Study the effect of Temperature on Salivary amylase act	tivity
12	Study the effect of substrate concentration on salivary a	mylase activity

BP204T	PATHOPHYSIOLOGY (THEORY)	45 Hours
Scope: Pat	hophysiology is the study of causes of diseases and reactions of	the body to such
disease producing causes. This course is designed to impart a thorough knowledge of the		
relevant aspects of pathology of various conditions with reference to its pharmacological		
applications, and understanding of basic pathophysiological mechanisms.		
	COURSE OUTCOMES	
CO 1	Understand the etiology, pathogenesis, and clinical manifestat diseases.	ions of common
CO 2	Explain the physiological and biochemical basis of disease proce	sses.
CO 3	Analyze the impact of diseases on various body systems.	
CO 4	Understand the principles of disease prevention and manageme	nt.
CO 5	Apply knowledge of pathophysiology in the context of drug the practice.	rapy and clinical
	COURSE CONTENT	
Unit I	Basic principles of Cell injury and Adaptation:	10 hours
	Introduction, definitions, Homeostasis, Components and Types	10 110413
	of Feedback systems, Causes of cellular injury, Pathogenesis	
	(Cell membrane damage, Mitochondrial damage, Ribosome	
	damage, nuclear damage), Morphology of cell injury – Adaptive	
	changes (Atrophy, Hypertrophy, hyperplasia, Metaplasia,	
	Dysplasia), Cell swelling, Intra cellular accumulation,	
	Calcification, Enzyme leakage and Cell Death Acidosis &	
	Alkalosis, Electrolyte imbalance	
	Basic mechanism involved in the process of inflammation	
	and repair: Introduction, Clinical signs of inflammation,	
	Different types of Inflammation, Mechanism of Inflammation -	
	Alteration in vascular permeability and blood flow, migration of	7
	WBC's, Mediators of inflammation, Basic principles of wound	
	healing in the skin, Pathophysiology of Atherosclerosis	
Unit II	Cardiovascular System: Hypertension, congestive heart	10
	failure, ischemic heart disease (angina, myocardial infarction,	
	atherosclerosis and arteriosclerosis)	hours
	Respiratory system: Asthma, Chronic obstructive airways	
	diseases.	
	Renal system: Acute and chronic renal failure.	
Unit III	Hematological Diseases:	10 hours
	Iron deficiency, megaloblastic anemia (Vit B12 and folic acid),	
	sickle cell anemia, thalassemia, hereditary acquired anemia	
	hemophilia	
	Endocrine system: Diabetes, thyroid diseases, disorders of sex	
	hormones	

	Nervous system: Epilepsy, Parkinson's disease, stroke,	
	psychiatric disorders: depression, schizophrenia and	
	Alzheimer's disease.	
	Gastrointestinal system: Peptic Ulcer	
Unit IV	Inflammatory bowel diseases, jaundice, hepatitis (A, B, C, D, E, F)	10 hours
	alcoholic liver disease.	
	Disease of bones and joints: Rheumatoid arthritis, osteoporosis	
	and gout	
	Principles of cancer: Classification, etiology and pathogenesis of	
	cancer	
	Diseases of bones and joints: Rheumatoid Arthritis,	
	Osteoporosis, Gout	
	Principles of Cancer: Classification, etiology and pathogenesis	
	of Cancer	
Unit V	Infectious diseases: Meningitis, Typhoid, Leprosy, Tuberculosis	09
	Urinary tract infections: Sexually transmitted diseases: AIDS,	hours
	Syphilis, Gonorrhea	

- 1. Vinay Kumar, Abul K. Abas, Jon C. Aster; Robbins & Cotran Pathologic Basis of Disease; South Asia edition; India; Elsevier; 2014.
- 2. Harsh Mohan; Text book of Pathology; 6 th edition; India; Jaypee Publications; 2010.
- 3. Laurence B, Bruce C, Bjorn K.; Goodman Gilman's The Pharmacological Basis of Therapeutics; 12th edition; New York; McGraw-Hill; 2011.
- 4. Best, Charles Herbert 1899-1978; Taylor, Norman Burke 1885-1972; West, John B (John Burnard); Best and Taylor's Physiological basis of medical practice; 12th ed; united states;
- 5. William and Wilkins, Baltimore;1991 [1990 printing].
- 6. Nicki R. Colledge, Brian R. Walker, Stuart H. Ralston; Davidson's Principles and Practice of Medicine; 21st edition; London; ELBS/Churchill Livingstone; 2010.
- 7. Guyton A, John. E Hall; Textbook of Medical Physiology; 12 th edition; WB Saunders Company; 2010.
- 8. Joseph DiPiro, Robert L. Talbert, Gary Yee, Barbara Wells, L. Michael Posey; Pharmacotherapy: A Pathophysiological Approach; 9 th edition; London; McGraw-Hill Medical; 2014.
- 9. V. Kumar, R. S. Cotran and S. L. Robbins; Basic Pathology; 6 th edition; Philadelphia; WB Saunders Company; 1997.
- 10. Roger Walker, Clive Edwards; Clinical Pharmacy and Therapeutics; 3 rd edition; London; Churchill Livingstone publication; 2003.

Recommended Journals

- 1. The Journal of Pathology. ISSN: 1096-9896 (Online)
- 2. The American Journal of Pathology. ISSN: 0002-9440
- 3. Pathology. 1465-3931 (Online)
- 4. International Journal of Physiology, Pathophysiology and Pharmacology. ISSN: 1944-8171 (Online)
- 5. Indian Journal of Pathology and Microbiology. ISSN-0377-4929.

BP205T	COMPUTER APPLICATIONS IN PHARMACY (Theory)	30 Hrs (2 Hrs/Week)
o mi :		

Scope: This subject deals with the introduction Database, Database Management system,

computer application in clinical studies and use of databases.		
COURSE OUTCOMES		
CO 1	Know the various types of computer applications used in the field of pharmacy.	
CO 2	Gain knowledge about different types of databases and database mana systems.	agement
CO 3	Apply databases effectively in various aspects of pharmacy, including studies and drug information management.	g clinical
CO 4	Understand web technologies, including HTML, XML, CSS, and servand their applications in pharmacy.	ver products,
CO 5	Use bioinformatics and data analysis tools in preclinical development, including chromatographic data analysis, Laboratory Information Management Systems (LIMS), and Text Information Management Systems (TIMS).	
COURSE CONTENT		
Unit I	Number system: Binary number system, Decimal number system, Octal number system, Hexadecimal number systems, conversion decimal to binary, binary to decimal, octal to binary etc, binary addition, binary subtraction – One's complement, Two's complement method, binary multiplication, binary division. Concept of Information Systems and Software: Information gathering, requirement and feasibility analysis, data flow diagrams, process specifications, input/output design, process life cycle, planning and managing the project	06 hours
Unit II	Web technologies: Introduction to HTML, XML, CSS and Programming languages, introduction to web servers and Server Products Introduction to databases, MYSQL, MS ACCESS, Pharmacy Drug database	06 hours
Unit III	Application of computers in Pharmacy – Drug information storage and retrieval, Pharmacokinetics, Mathematical model in Drug design, Hospital and Clinical Pharmacy, Electronic Prescribing and discharge (EP) systems, barcode medicine identification and automated dispensing of drugs, mobile technology and adherence monitoring Diagnostic System, Labdiagnostic System, Patient Monitoring System, Pharma	06 hours

	Information System	
Unit IV	Bioinformatics: Introduction, Objective of Bioinformatics,	06 hours
	Bioinformatics Databases, Concept of Bioinformatics, Impact of	
	Bioinformatics in Vaccine Discovery	
Unit V	Computers as data analysis in Preclinical development:	06 hours
	Chromatographic dada analysis (CDS), Laboratory Information	
	management System (LIMS) and Text Information Management	
	System (TIMS)	

- 1. Computer Application in Pharmacy William E. Fassett –Lea and Febiger, 600 South Washington Square, USA, (215) 922-1330.
- 2. Computer Application in Pharmaceutical Research and Development -Sean Ekins -Wiley-Interscience, A John Willey and Sons, INC., Publication, USA
- 3. Bioinformatics (Concept, Skills and Applications) S.C. Rastogi-CBS Publishers and Distributors, 4596/1- A, 11 Darya Gani, New Delhi – 110 002(INDIA)
- 4. Microsoft office Access 2003, Application Development Using VBA, SQL Server, DAP and Infopath - Cary N. Prague - Wiley Dreamtech India (P) Ltd., 4435/7, Ansari Road, Daryagani, New Delhi - 110002

BP210P	COMPUTER APPLICATIONS IN PHARMACY (Practical)	2 Hours/Week	
S.No.	Experiment		
1	Design a questionnaire using a word processing package to gather information about a particular disease.		
2	Create a HTML web page to show personal information.		
3	Retrieve the information of a drug and its adverse effects using online tools.		
4	Creating mailing labels using Label Wizard, generating label in MS WORD.		
5	Create a database in MS Access to store the patient information with the required fields using Access.		
6	Design a form in MS Access to view, add, delete, and modify the patient record in the database.		
7	Generating report and printing the report from patient database.		
8	Creating invoice table using MS Access.		
9	Drug information storage and retrieval using MS Access.		
10	Creating and working with queries in MS Access.		
11	Exporting Tables, Queries, Forms, and Reports to web pages.		
12	Exporting Tables, Queries, Forms, and Reports to XML pages.		

Scope: Environmental Sciences is the scientific study of the environmental system and the status of its inherent or induced changes on organisms. It includes not only the study of physical and biological characters of the environment but also the social and cultural factors and the impact of man on environment.

man on environment.				
COURSE OUTCOMES				
	Understand the various resources of environment and apply	_		
CO 1	environmental science to promote sustainable practices in pharmace	eutical industries		
	and healthcare.			
CO 2	Understand the principles of ecology and environmental science.			
CO 3	Analyze the effects of pollution on health and the environment.			
COURSE CONTENT				
Unit I	The Multidisciplinary nature of environmental studies Natural	10 hours		
	Resources	10 110413		
	Renewable and non-renewable resources:			
	Natural resources and associated problems			
	a) Forest resources; b) Water resources; c) Mineral resources;			
	d) Food resources; e) Energy resources; f) Land resources: Role			
	of an individual in conservation of natural resources.			
Unit II	Ecosystems	10 hours		
	Concept of an ecosystem.	10 110 1115		
	Structure and function of an ecosystem.			
	• Introduction, types, characteristic features, structure and			
	function of the ecosystems: Forest ecosystem; Grassland			
	ecosystem; Desert ecosystem; Aquatic ecosystems (ponds,			
	streams, lakes, rivers, oceans, estuaries)			
Unit III	Environmental Pollution: Air pollution; Water pollution; Soil	10 hours		
	pollution	10 110413		

- 1. Y.K. Sing, Environmental Science, New Age International Pvt, Publishers, Bangalore
- 2. Agarwal, K.C. 2001 Environmental Biology, Nidi Publ. Ltd. Bikaner.
- 3. Bharucha Erach, The Biodiversity of India, Mapin Pu blishing Pvt. Ltd., Ahmedabad 380 013, India,
- 4. Brunner R.C., 1989, Hazardous Waste Incineration, McGraw Hill Inc. 480p
- 5. Clark R.S., Marine Pollution, Clanderson Press Oxford
- 6. De A.K., Environmental Chemistry, Wiley Eastern Ltd.
- 7. Down of Earth, Centre for Science and Environment

B. PHARM (SEMESTER – III)			
BP301T	PHARMACEUTICAL ORGANIC CHEMISTRY -II (Theory)	45 Hours	
functions o	s subject is designed to impart fundamental knowledge on the structure of the various systems of the human body. It also helps in understance mechanisms. The subject provides the basic knowledge real the various disciplines of pharmacy.	nding both	
	COURSE OUTCOMES		
CO 1	Understand the structural features of benzene and its derivatives aromaticity, resonance, and substitution reactions, and apply this kn predict reactivity and orientation in electrophilic substitution reactions	owledge to	
CO 2	Understand and apply analytical constants to characterize fats and oils acid value, saponification value, and iodine value.	s, including	
CO 3	Valuate the structural and synthesis features contributing to their properties and apply this knowledge to pharmaceutical applications.		
CO 4	Analyze the acidity of phenols and aromatic acids and evaluate the impact of substituents on their acidity. Apply qualitative tests to identify phenols and assess the structure-activity relationships for various phenolic compounds.		
CO 5	Analyze how different groups affect the behavior of organic compounds, especially in aromatic and cycloalkane molecules.		
COURSE CONTENT			
be explain	ethods of preparation and reactions of compounds superscripted with ast ned. To emphasize on definition, types, classification, principles/mas, examples and difference		
Unit I	 Benzene and its derivatives a. Analytical, synthetic and other evidences in the derivation of structure of benzene, Orbital picture, resonance in benzene, aromatic characters, Huckel's rule b. Reactions of benzene - nitration, sulphonation, halogenation-reactivity, Friedelcrafts alkylation- reactivity, limitations, Friedelcrafts acylation. c. Substituents, effect of substituents on reactivity and orientation of mono substituted benzene compounds towards electrophilic substitution reaction d. Structure and uses of DDT, Saccharin, BHC and Chloramine 	10 hours	
Unit II	Phenols* - Acidity of phenols, effect of substituents on acidity, qualitative tests, Structure and uses of phenol, cresols, resorcinol,	10 hours	

	naphthols	
	Aromatic Amines* - Basicity of amines, effect of substituents on	
	basicity, and synthetic uses of aryl diazonium salts	
	Aromatic Acids* -Acidity, effect of substituents on acidity and	
	important reactions of benzoic acid.	
Unit III	Fats and Oils	10 hours
	a. Fatty acids – reactions.	To nours
	b. Hydrolysis, Hydrogenation, Saponification and Rancidity of oils,	
	Drying oils.	
	c. Analytical constants – Acid value, Saponification value, Ester	
	value, Iodine value, Acetyl value, Reichert Meissl (RM) value –	
	significance and principle involved in their determination.	
Unit IV	Polynuclear hydrocarbons:	001
	a. Synthesis, reactions	08 hours
	b. Structure and medicinal uses of Naphthalene, Phenanthrene,	
	Anthracene, Diphenylmethane, Triphenylmethane and their	
	derivatives	
Unit V	Cyclo alkanes*	0.5.1
	Stabilities – Baeyer's strain theory, limitation of Baeyer's strain	07 hours
	theory, Coulson and Moffitt's modification, Sachse Mohr's theory	
	(Theory of strainless rings), reactions of cyclopropane and	
	cyclobutane-only	
	cyclobatane only	
	RECOMMENDED BOOKS (LATEST EDITIONS)	
1.	Organic Chemistry by Morrison and Boyd	
2.	Organic Chemistry by I.L. Finar, Volume-I	
3.	Textbook of Organic Chemistry by B.S. Bahl & Arun Bahl.	
4.	Organic Chemistry by P.L. Soni	
5.	Practical Organic Chemistry by Mann and Saunders.	
6.	Vogel's text book of Practical Organic Chemistry	
7.	Advanced Practical organic chemistry by N.K. Vishnoi.	
8.	Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz	Z.

BP305P	PHARMACEUTICAL ORGANIC CHEMISTRY -II (Practical)	4 Hrs/Week	
Section	Experiments		
I	Experiments involving laboratory techniques		
	- Recrystallization		
	- Steam distillation		
II	Determination of following oil values (including standardizati	on of reagents)	
	- Acid value		
	- Saponification value		
	- Iodine value		
III	Preparation of compounds		
	- Benzanilide/Phenyl benzoate/Acetanilide from Aniline/Phenol/Aniline by		
	acylation reaction		
	- 2,4,6-Tribromo aniline/Para bromo acetanilide from Aniline/Acetanilide by		
	halogenation (Bromination) reaction		
	- 5-Nitro salicylic acid/Meta di nitro benzene from Salicylic acid/Nitro benzene		
	by nitration reaction		
	- Benzoic acid from Benzyl chloride by oxidation reaction		
	- Benzoic acid/Salicylic acid from alkyl benzoate/alkyl salicylate	by hydrolysis	
	reaction		
	- 1-Phenyl azo-2-napthol from Aniline by diazotization and coupl	ing reactions	
	- Benzil from Benzoin by oxidation reaction		
	- Dibenzal acetone from Benzaldehyde by Claisen-Schmidt reacti	on	
	- Cinnamic acid from Benzaldehyde by Perkin reaction		
	- p-Iodo benzoic acid from p-Amino benzoic acid		

BP302T	PHYSICAL PHARMACEUTICS-I (Theory)	45 Hours	
Scope: The course deals with the various physical and physicochemical properties, and principles			
	involved in dosage forms/formulations. Theory and practical components of the subject help the		
_	get a better insight into various areas of formulation research and de	evelopment, and	
stability stu	dies of pharmaceutical dosage forms.		
	COURSE OUTCOMES		
CO 1	Understand and apply the principles of drug solubility and the factor	rs affecting it.	
CO 2	Describe and analyze the states and physicochemical properties of n pharmaceuticals.	natter relevant to	
CO 3	Understand and explain surface and interfacial phenomena, includi techniques and applications in pharmaceuticals.		
CO 4	Understand the principles of complexation and protein binding, their and their significance in drug action and stability.		
CO 5	Apply knowledge of pH, buffer systems, and isotonic solutions in formulations and biological systems.	n pharmaceutical	
	COURSE CONTENT		
Unit I	Solubility of drugs: Solubility expressions, mechanisms of solute solvent interactions, ideal solubility parameters solvation & association, quantitative approach to the factors influencing solubility of drugs, diffusion principles in biological systems. Solubility of gas in liquids, solubility of liquids in liquids, (Binary solutions, ideal solutions) Raoult's law, real solutions. Partially miscible liquids, Critical solution temperature and applications. Distribution law, its limitations and applications States of Matter and properties of matter: State of matter changes in the state of matter, latent heats, vapour pressure sublimation critical point, eutectic mixtures, gases, aerosols—inhalers, relative humidity, liquid complexes, liquid crystals glassy states, solid- crystalline, amorphous & polymorphism Physicochemical properties of drug molecules: Refractive	10 hours	
Unit III	index, optical rotation, dielectric constant, dipole moment dissociation constant, determinations and applications Surface and interfacial phenomenon: Liquid interface, surface & interfacial tensions, surface free energy, measurement of surface & interfacial tensions, spreading coefficient, adsorption at liquid interfaces, surface active agents, HLB Scale, solubilisation detergency, adsorption at solid interface.	08 hours	
Unit IV	Complexation and protein binding: Introduction, Classification o	of 08 hours	

	Complexation, Applications, methods of analysis, protein binding, Complexation and drug action, crystalline structures of complexes and thermodynamic treatment of stability constants.	
Unit V	pH, buffers and Isotonic solutions: Sorensen's pH scale, pH determination (electrometric and calorimetric), applications of buffers, buffer equation, buffer capacity, buffers in pharmaceutical and biological systems, buffered isotonic solutions.	07 hours

- 1. Physical Pharmacy by Alfred Martin
- 2. Experimental Pharmaceutics by Eugene, Parott.
- 3. Tutorial Pharmacy by Cooper and Gunn.
- 4. Stocklosam J. Pharmaceutical Calculations, Lea & Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to MarcelDekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical Dosage forms. Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc. 7. Physical Pharmaceutics by Ramasamy C and ManavalanR.
- 7. Laboratory Manual of Physical Pharmaceutics, C.V.S. Subramanyam, J. Thimma settee
- 8. Physical Pharmaceutics by C.V.S. Subramanyam
- 9. Test book of Physical Pharmacy, by Gaurav Jain & Roop K. Khar

BP306P	PHYSICAL PHARMACEUTICS - I (Practical)	4 Hours/Week	
S.No.	Experiment		
1	Determination of the solubility of a drug at room temperature	re	
2	Determination of pKa value by Half Neutralization/Henderse equation	on-Hasselbalch	
3	Determination of partition coefficient of benzoic acid in benz	zene and water	
4	Determination of partition coefficient of iodine in CCl ₄ and water		
5	Determination of % composition of NaCl in a solution using phenol-water system		
	by CST method		
6	Determination of surface tension of given liquids by drop count and drop weight method		
7	Determination of HLB number of a surfactant by saponificati	ion method	
8	Determination of Freundlich and Langmuir constants using activated charcoal		
9	Determination of critical micellar concentration of surfactan	ts	
10	Determination of stability constant and donor-acceptor ratio	of PABA-Caffeine	
	complex by solubility method		
11	Determination of stability constant and donor-acceptor ratio	of Cupric-Glycine	
	complex by pH titration method		

BP303T	PHARMACEUTICAL MICROBIOLOGY (Theory)	45 Hours	
Scope: Study of all categories of microorganisms especially for the production of alcohol antibiotics, vaccines, vitamins enzymes etc			
COURSE OUTCOMES			

	COURSE OUTCOMES	
CO 1	Understand the fundamentals of microbiology relevant to pharmaceutical sciences, including microbial structure, growth, and metabolism.	
CO 2	Analyze the role of microorganisms in pharmaceutical manufacturing, control, and product stability assessment.	contamination
CO 3	Discuss the principles and techniques of sterilization, disinfection, enumeration in pharmaceutical quality assurance.	and microbial
CO 4	Evaluate the microbiological quality of pharmaceutical raw mater products, and manufacturing environments.	erials, finished
CO 5	Apply microbiological concepts and techniques in the development are aseptic processing and microbial control strategies.	nd validation of
	COURSE CONTENT	
Unit I	its importance. Introduction to Prokaryotes and Eukaryotes Study of ultra-structure and morphological classification of bacteria, nutritional requirements, raw materials used for culture media and physical parameters for growth, growth curve, isolation and preservation methods for pure cultures, cultivation of anaerobes, quantitative measurement of bacterial growth (total & viable count). Study of different types of phase contrast microscopy, dark field microscopy and electron microscopy.	10 hours
Onica	Gram's &Acid-fast staining) and biochemical tests (IMViC). Study of principle, procedure, merits, demerits and applications of physical, chemical gaseous, radiation and mechanical method of sterilization. Evaluation of the efficiency of sterilization methods. Equipment's employed in large scale sterilization. Sterility indicators.	10 hours
Unit II	Study of morphology, classification, reproduction/replication and cultivation of Fungi and Viruses. Classification and mode of action of disinfectants Factors influencing disinfection, antiseptics and their evaluation. For bacteriostatic and bactericidal actions Evaluation of bactericidal & Bacteriostatic. Sterility testing of products (solids, liquids, ophthalmic and other sterile products) according to IP, BP and USP.	10 hours

Unit IV	Designing of aseptic area, laminar flow equipment's; study of different sources of contamination in an aseptic area and methods of prevention, clean area classification. Principles and methods of different microbiological assay. Methods for standardization of antibiotics, vitamins and amino acids. Assessment of a new antibiotic.	08 hours
Unit V	Types of spoilage, factors affecting the microbial spoilage of pharmaceutical products, sources and types of microbial contaminants, assessment of microbial contamination and spoilage. Preservation of pharmaceutical products using antimicrobial agents, evaluation of microbial stability of formulations. Growth of animal cells in culture, general procedure for cell culture, Primary, established and transformed cell cultures. Application of cell cultures in pharmaceutical industry and research.	07 hours

- 1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.
- 2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.
- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. I.P., B.P., U.S.P.- latest editions.
- 10. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 11. Edward: Fundamentals of Microbiology.
- 12. N.K. Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 13. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company

BP307P	PHARMACEUTICAL MICROBIOLOGY (Practical)	4 Hours/Week
S.No.	Experiment	
1	Introduction and study of different equipment and processing	g, e.g., B.O.D. incubator,
	laminar flow, aseptic hood, autoclave, hot air sterilizer, deep	freezer, refrigerator,
	microscopes used in experimental microbiology.	
2	Sterilization of glassware, preparation, and sterilization of n	nedia.
3	Subculturing of bacteria and fungus. Nutrient stabs and slants preparations.	
4	Staining methods - Simple, Gram's staining, and acid-fast staining (Demonstration	
	with practical).	
5	Isolation of pure culture of micro-organisms by multiple streak plate technique and	
	other techniques.	
6	Microbiological assay of antibiotics by cup plate method an	d other methods.
7	Motility determination by hanging drop method.	
8	Sterility testing of pharmaceuticals.	
9	Bacteriological analysis of water.	
10	Biochemical tests.	

		45 He	
BP304T	PHARMACEUTICAL ENGINEERING (THEORY)	45 Hours	
_	Scope: This course is designed to impart a fundamental knowledge on the art and science		
of various ι	unit operations used in pharmaceutical industry.		
	COURSE OUTCOMES		
CO 1	Understand fluid flow principles, size reduction mechanisms, and size separation		
	techniques used in pharmaceutical processes.	and distillation	
CO 2	Gain knowledge of heat transfer methods, evaporation processes, techniques essential for pharmaceutical manufacturing.	and distillation	
CO 3	Comprehend drying and mixing principles and their applications in production.	pharmaceutical	
CO 4	Understand filtration and centrifugation principles and their roles in manufacturing.	pharmaceutical	
CO 5	Learn about materials selection for pharmaceutical plant constru prevention methods, and basics of material handling systems.	ction, corrosion	
	COURSE CONTENT		
Unit I	Flow of fluids: Types of manamatana Daymalda nymbay and ita		
Uniti	Flow of fluids: Types of manometers, Reynolds number and its significance, Bernoulli's theorem and its applications, Energy	10 hours	
	losses, Orifice meter, Venturimeter, Pitot tube and Rotameter.		
	Size Reduction: Objectives, Mechanisms & Laws governing size		
	reduction, factors affecting size reduction, principles,		
	construction, working, uses, merits and demerits of Hammer		
	mill, ball mill, fluid energy mill, Edge runner mill & end runner mill.		
	Size Separation: Objectives, applications & mechanism of size		
	separation, official standards of powders, sieves, size		
	separation Principles, construction, working, uses, merits and		
	demerits of Sieve shaker, cyclone separator, Air separator, Bag		
	filter & elutriation tank.		
Unit II	Heat Transfer: Objectives, applications & Heat transfer	10	
	mechanisms. Fourier's law, Heat transfer by conduction, convection	hours	
	& radiation. Heat interchangers & heat exchangers. 82 •	nours	
	Evaporation: Objectives, applications and factors influencing		
	evaporation, differences between evaporation and other heat process. principles, construction, working, uses, merits and demerits		
	of Steam jacketed kettle, horizontal tube evaporator, climbing film		
	evaporator, forced circulation evaporator, multiple effect		
	evaporator& Economy of multiple effect evaporator.		
	Distillation: Basic Principles and methodology of simple		
	distillation, flash distillation, fractional distillation, distillation		
	under reduced pressure, steam distillation & molecular distillation		
Unit III	Drying: Objectives, applications & mechanism of drying process,	08 hours	

	measurements & applications of Equilibrium Moisture content, rate		
	of drying curve. principles, construction, working, uses, merits and		
	demerits of Tray dryer, drum dryer spray dryer, fluidized bed dryer,		
	vacuum dryer, freeze dryer.		
	Mixing: Objectives, applications & factors affecting mixing,		
	Difference between solid and liquid mixing, mechanism of solid		
	mixing, liquids mixing and semisolids mixing. Principles,		
	Construction, Working, uses, Merits and Demerits of Double cone		
	blender, twin shell blender, ribbon blender, Sigma blade mixer,		
	planetary mixers, Propellers, Turbines, Paddles & Silverson		
	Emulsifier.		
Unit IV	Filtration: Objectives, applications, Theories & Factors influencing	08 hours	
	filtration, filter aids, filter medias. Principle, Construction, Working,		
	Uses, Merits and demerits of plate & frame filter, filter leaf, rotary		
	drum filter, Meta filter & Cartridge filter, membrane filters and Seidtz		
	filter.		
	Centrifugation: Objectives, principle & applications of		
	Centrifugation, principles, construction, working, uses, merits and		
	demerits of Perforated basket centrifuge, non-perforated basket		
	centrifuge, semi continuous centrifuge & super centrifuge.		
Unit V	Materials of pharmaceutical plant construction, Corrosion and	07 hours	
	its prevention: Factors affecting during materials selected for		
	pharmaceutical plant construction, Theories of corrosion, types of		
	corrosion and their prevention. Ferrous and nonferrous metals,		
	inorganic and organic nonmetals, basic of material handling systems.		
DECOMMENDED DOOKS (LATEST EDITIONS)			

- 1. Introduction to chemical engineering Walter L Badger & Julius Banchero, Latest
- 2. Solid phase extraction, Principles, techniques and applications by Nigel J.K. Simpson- Latest edition.
- 3. Unit operation of chemical engineering Mcabe Smith, Latest edition.
- 4. Pharmaceutical engineering principles and practices C.V.S Subrahmanyam et al., Latest
- 5. edition.
- 6. Remington practice of pharmacy- Martin, Latest edition.
- 7. Theory and practice of industrial pharmacy by Lachmann., Latest edition.
- 8. Physical pharmaceutics- C.V.S Subrahmanyam et al., Latest edition.
- 9. Cooper and Gunn's Tutorial pharmacy, S.J. Carter, Latest edition.

BP308P	PHARMACEUTICAL ENGINEERING (Practical)	4 Hours/Week	
S.No.	Experiment		
I	Determination of radiation constant of brass, iron, unp	ainted, and painted glass.	
II	Steam distillation - Calculation of the efficiency of ste	am distillation.	
III	Determination of the overall heat transfer coefficient b	y heat exchanger.	
IV	Construction of drying curves (for calcium carbonate a	and starch).	
V	Determination of moisture content and loss on drying.		
VI	Determination of humidity of air using wet and dry bu	lb temperatures, and Dew	
	point method.		
VII	Description of construction, working, and application	of Pharmaceutical Machinery	
	(e.g., rotary tablet machine, fluidized bed coater, fluid energy mill, dehumidifier).		
VIII	Size analysis by sieving – Evaluation of size distribution of tablet granulations,		
	construction of size frequency curves including arithmetic and logarithmic		
	probability plots.		
IX	Size reduction: Verification of size reduction laws using ball mill, determination of		
	Kicks, Rittinger's, Bond's coefficients, power requirer	ment, and critical speed of Ball	
	Mill.		
X	Demonstration of colloid mill, planetary mixer, fluidized bed dryer, freeze dryer, and		
	other major equipment.		
XI	Factors affecting Rate of Filtration and Evaporation (Surface area, Concentration,		
	Thickness/viscosity).		
XII	Study of the effect of time on the Rate of Crystallization.		
XIII	Calculation of the uniformity index for a given sample using Double Cone Blender.		

B. PHARM (SEMESTER – IV)			
BP401T	PHARMACEUTICAL ORGANIC CHEMISTRY -III (Theory)	45 Hours	
and organic	Scope: This subject imparts knowledge on stereo-chemical aspects of organic compounds and organic reactions, important named reactions, chemistry of important hetero cyclic compounds. It also emphasizes on medicinal and other uses of organic compounds.		
	COURSE OUTCOMES		
CO 1	Demonstrate a thorough understanding of stereochemical aspects compounds, including optical isomerism, geometrical isome conformational isomerism, enabling the interpretation of molecular st behaviour.	erism, and	
CO 2	Analyse the mechanisms and stereochemistry of organic reactions releasynthesis of medicinal compounds.		
CO 3	Discuss the principles of retrosynthetic analysis and multi-step synthes for complex organic molecules.		
CO 4	Evaluate the medicinal uses and other applications of organic compounds, with a focus on stereochemistry, recognizing the importance of chirality and spatial arrangement in drug design and other fields.		
CO 5	Apply knowledge of named reactions, including metal hydride reductions, oxidations, rearrangements, and other synthetic transformations, to design efficient synthetic routes for the preparation of complex organic molecules.		
	COURSE CONTENT		
Note: To e	mphasize on definition, types, mechanisms, examples, uses/applicati	ons	
Unit I	Stereo isomerism Optical isomerism – Optical activity, enantiomerism, diastereoisomerism, meso compounds Elements of symmetry, chiral and achiral molecules, DL system of nomenclature of optical isomers, sequence rules, RS system of nomenclature of optical isomers, Reactions of chiral molecules, Racemic modification and resolution of racemic mixture, Asymmetric synthesis: partial and absolute.	10 hours	
Unit II	Geometrical isomerism Nomenclature of geometrical isomers (Cis Trans, EZ, Syn Anti systems) Methods of determination of configuration of geometrical isomers. Conformational isomerism in Ethane, n-Butane and Cyclohexane. Stereo isomerism in biphenyl compounds (Atropoisomerism) and	10 hours	

	conditions for optical activity.	
	Stereospecific and stereoselective reactions	
Unit III	Heterocyclic compounds:	10 hours
	Nomenclature and classification, Synthesis, reactions and	10 Hours
	medicinal uses of following compounds/derivatives Pyrrole,	
	Furan, and Thiophene	
Unit IV	Synthesis, reactions and medicinal uses of following	08 hours
	compounds/derivatives. Pyrazole, Imidazole, Oxazole and	
	Thiazole. Pyridine, Quinoline, Isoquinoline, Acridine and Indole.	
	Basicity of pyridine Synthesis and medicinal uses of Pyrimidine,	
	Purine, azepines, and their derivatives.	
Unit V	Reactions of synthetic importance	
	Metal hydride reduction (NaBH4 and LiAlH4), Clemmensen	07 hours
reduction, Birch reduction, Wolff Kishner reduction. Oppenauer-		
oxidation and Dakin reaction. Beckmanns rearrangement and		
	Schmidt rearrangement. Claisen-Schmidt condensation	
	RECOMMENDED BOOKS (LATEST EDITIONS)	
1.	Organic chemistry by I.L. Finar, Volume-I & II.	
2.	A text book of organic chemistry – Arun Bahl, B.S. Bahl.	
	Heterocyclic Chemistry by Raj K. Bansal	
	Organic Chemistry by Morrison and Boyd	
5.	Heterocyclic Chemistry by T.L. Gilchrist	

BP402T	MEDICINAL CHEMISTRY – I (Theory)	45 Hours	
Scope: This subject is designed to impart fundamental knowledge on the structure, chemistry, and			
therapeutic	therapeutic value of drugs. The subject emphasizes on structure-activity relationships of drugs, the		
	of physicochemical properties and the metabolism of drugs. The	ne syllabus also	
emphasizes	on chemical synthesis of important drugs under each class.		
	COURSE OUTCOMES		
CO 1	Demonstrate proficiency in writing the chemical synthesis of selected principles of organic chemistry to prepare pharmaceutical compound safely.		
CO 2	Analyze the chemistry of drugs in relation to their pharmacological acunderstanding the mechanisms of action and the influence of properties on drug behavior.	•	
CO 3	Evaluate drug metabolic pathways, recognizing potential adverse effects and assessing the therapeutic value of drugs in clinical settings, and understand the principles of drug metabolism, including Phase I and Phase II reactions.		
CO 4	Evaluate the structure-activity relationships (SAR) of drugs as	nd predict their	
	pharmacological profiles using computational methods.		
CO 5	Apply acquired knowledge to critically evaluate the development, classification, and uses of drugs covered in the course, integrating concepts of chemistry and pharmacology to understand their therapeutic applications.		
	COURSE CONTENT		
action, use	he development of the following classes of drugs, Classification, es of drugs mentioned in the course, Structure-activity relationsh	ip of a selective	
	ugs as specified in the course and synthesis of drugs superscripte	ed (*)	
Unit I	Introduction to Medicinal Chemistry	10 hours	
	History and development of medicinal chemistry		
	Physicochemical properties in relation to biological action Ionization, Solubility, Partition Coefficient, Hydrogen bonding		
	Protein binding, Chelation, Bioisosterism, Optical and		
	Geometrical isomerism.	4	
	Drug metabolism		
	Drug metabolism principles- Phase I and Phase II.		
	Factors affecting drug metabolism including stereochemical	l	
	aspects.		
Unit II	Drugs acting on Autonomic Nervous System	10 hours	
	Adrenergic Neurotransmitters:		
	Biosynthesis and catabolism of catecholamine.		
	Adrenergic receptors (Alpha & Beta) and their distribution.		

	Sympathomimetic agents: SAR of Sympathomimetic agents	
	Direct acting: Nor-epinephrine, Epinephrine, Phenylephrine*,	
	Dopamine, Methyldopa, Clonidine, Dobutamine, Isoproterenol,	
	Terbutaline, Salbutamol*, Bitolterol,	
	Naphazoline, Oxymetazoline and Xylometazoline.	
	Indirect-acting agents: Hydroxy amphetamine,	
	Pseudoephedrine, Propylhexedrine.	
	Agents with mixed mechanism: Ephedrine, Metaraminol.	
	Adrenergic Antagonists:	
	Alpha-adrenergic blockers: Tolazoline*, Phentolamine,	
	Phenoxybenzamine, Prazosin, Dihydroergotamine,	
	Methysergide.	
	Beta-adrenergic blockers: SAR of beta blockers, Propranolol*,	
	Metipranolol, Atenolol, Betazolol, Bisoprolol, Esmolol,	
	Metoprolol, Labetalol, Carvedilol.	
Unit III	Cholinergic neurotransmitters:	10 h
	Biosynthesis and catabolism of acetylcholine.	10 hours
	Cholinergic receptors (Muscarinic & Nicotinic) and their	
	distribution.	
	Parasympathomimetic agents: SAR of Parasympathomimetic	
	agents	
	Direct-acting agents: Acetylcholine, Carbachol*, Bethanechol,	
	Methacholine, Pilocarpine.	
	Indirect acting/ Cholinesterase inhibitors (Reversible &	
	Irreversible): Physostigmine, Neostigmine*, Pyridostigmine,	
	Edrophonium chloride, Tacrine hydrochloride, Ambenonium	
	chloride, Isofluorphate, Echothiopate iodide, Parathion, Malathion.	
	Cholinesterase reactivator: Pralidoxime chloride.	
	Cholinergic Blocking agents: SAR of cholinolytic agents	
	Solanaceous alkaloids and analogs: Atropine sulfate, Hyoscyamine	
	sulphate, Scopolamine hydrobromide, Homatropine hydrobromide,	
	Ipratropium bromide*.	
	Synthetic cholinergic blocking agents: Tropicamide,	
	Cyclopentolate hydrochloride, Clidinium bromide, Dicyclomine	
	hydrochloride*, Glycopyrrolate, Methantheline bromide,	
	Propantheline bromide, Benztropine mesylate, Orphenadrine citrate,	
	Biperiden hydrochloride, Procyclidine hydrochloride*,	
	Tridihexethyl chloride, Isopropamide iodide, Ethopropazine	
	hydrochloride.	
Unit IV	Drugs acting on Central Nervous System	08 hours
Omtiv	A. Sedatives and Hypnotics:	oo nours
	Benzodiazepines: SAR of Benzodiazepines, Chlordiazepoxide,	
	Diazepam*, Oxazepam, Clorazepate, Lorazepam, Alprazolam,	
	Diazepani, Oxazepani, Ciorazepane, Lorazepani, Aiprazolam,	

Zolpidem

Barbiturates: SAR of barbiturates, Barbital*, Phenobarbital, Mephobarbital, Amobarbital, Butobarbital, Pentobarbital, Secobarbital

Miscellaneous:

Amides & imides: Glutethimide.

Alcohol & their carbamate derivatives: Meprobamate, Ethchlorvynol. Aldehyde & their derivatives: Triclofos sodium, Paraldehyde.

B. Antipsychotics

Phenothiazines: SAR of Phenothiazines - Promazine hydrochloride, Chlorpromazine hydrochloride*, Triflupromazine, Thioridazine hydrochloride, Piperacetazine hydrochloride, Prochlorperazine maleate, Triflupromazine hydrochloride.

Ring Analogues of Phenothiazines: Chlorprothixene, Thiothixene, Loxapine succinate, Clozapine.

Fluro butyrophenones: Haloperidol, Droperidol, Risperidone.

Beta amino ketones: Molindone hydrochloride.

Benzamides: Sulpieride.

C.Anticonvulsants: SAR of Anticonvulsants, mechanism of

anticonvulsant action

Barbiturates: Phenobarbitone, Methabarbital.

Hydantoins: Phenytoin*, Mephenytoin, Ethotoin

Oxazolidine diones: Trimethadione, Paramethadione

Succinimides: Phensuximide, Methsuximide, Ethosuximide* **Urea and monoacylureas:** Phenacemide, Carbamazepine*

Benzodiazepines: Clonazepam

Miscellaneous: Primidone, Valproic acid, Gabapentin, Felbamate

Unit V Drugs

Drugs acting on Central Nervous System

General anesthetics:

Inhalation anesthetics: Halothane*, Methoxyflurane, Enflurane,

Sevoflurane, Isoflurane, Desflurane.

Ultra short-acting barbiturates: Methohexital sodium*,

Thiamylal sodium, Thiopental sodium.

Dissociative anesthetics: Ketamine hydrochloride. *

Narcotic and non-narcotic analgesics

Morphine and related drugs: SAR of Morphine analogs, Morphine sulphate, Codeine, Meperidine hydrochloride, Anilerdine hydrochloride, Diphenoxylate hydrochloride, Loperamide hydrochloride, Fentanyl citrate*, Methadone hydrochloride*, Propoxyphene hydrochloride, Pentazocine, Levorphanol tartarate.

Narcotic antagonists: Nalorphine hydrochloride, Levallorphan

07 hours

tartarate, Naloxone hydrochloride.	
Anti-inflammatory agents: Sodium salicylate, Aspirin,	
Mefenamic acid*, Meclofenamate, Indomethacin, Sulindac,	
Tolmetin, Zomepirac, Diclofenac, Ketorolac, Ibuprofen*,	
Naproxen, Piroxicam, Phenacetin, Acetaminophen, Antipyrine,	
Phenylbutazone.	

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I. Vogel.

BP406P	MEDICINAL CHEMISTRY – I (Practical) 4 Hours/Wee	ek
Category	Experiment	
I.	Preparation of Drugs/Intermediates	
	1. 1,3-pyrazole	
	2. 1,3-oxazole	
	3. Benzimidazole	
	4. Benztriazole	
	5. 2,3-diphenyl quinoxaline	
	6. Benzocaine	
	7. Phenytoin	
	8. Phenothiazine	
	9. Barbiturate	
II.	Assay of Drugs	
	1. Chlorpromazine	
	2. Phenobarbitone	
	3. Atropine	
	4. Ibuprofen	
	5. Aspirin	
	6. Furosemide	
III.	Determination of Partition Coefficient for Any Two Drugs	

BP403T	PHYSICAL PHARMACEUTICS-II (Theory)	45 Hours	
Scope: The course deals with the various physica and physicochemical properties, and principles			
	involved in dosage forms/formulations. Theory and practical components of the subject help the		
student to g	get a better insight into various areas of formulation research and de	evelopment, and	
stability stu	dies of pharmaceutical dosage forms.		
	COURSE OUTCOMES		
CO 1	Understand the classification and properties of colloidal dispersions	s, including their	
CO 2	stability influenced by electrolytes and protective agents.		
CO 2	Gain knowledge of rheological principles it's applications on pharm		
CO 3	Learn about the formulation principles and properties of emulsions a	and suspensions.	
CO 4	Understand the fundamental and derived properties of particles.	C 4 1 '1' 4'	
CO F	Learn reaction kinetics, factors affecting drug degradation, methods		
CO 5	against common reactions, accelerated stability testing, and prevent	ion of photolytic	
	degradation in drug stability.		
	COURSE CONTENT		
Unit I	Colloidal dispersions: Classification of dispersed systems &	07 hours	
	their general characteristics, size & shapes of colloidal particles	, O7 Hours	
	classification of colloids &comparative account of their general	l	
	properties. Optical, kinetic & electrical properties. Effect of	f	
	electrolytes, coacervation, peptization& protective action.		
Unit II	Rheology: Newtonian systems, law of flow, kinematic viscosity	10 hours	
	effect of temperature, non-Newtonian systems, pseudoplastic	, 10 110 113	
	dilatant, plastic, thixotropy, thixotropy in formulation	,	
	determination of viscosity, capillary, falling Sphere, rotational	l	
	viscometers.		
	Deformation of solids: Plastic and elastic deformation, Heckel		
	equation, Stress, Strain, Elastic Modulus		
Unit III	Coarse dispersion: Suspension, interfacial properties of suspended	10 Hours	
	particles, settling in suspensions, formulation of flocculated and		
	deflocculated suspensions. Emulsions and theories of emulsification		
	microemulsion and multiple emulsions; Stability of emulsions		
	preservation of emulsions, rheological properties of emulsions and	d	
II!A II7	emulsion formulation by HLB method.	10 h a	
Unit IV	Micromeretics: Particle size and distribution, mean particle size		
	number and weight distribution, particle number, methods for		
	determining particle size by different methods, counting and separation method, particle shape specific surface methods to		
	separation method, particle shape, specific surface, methods fo determining surface area, permeability, adsorption, derived		
	properties of powders, porosity, packing arrangement, densities		
	bulkiness & flow properties.	''	
	ountilless & non properties.		

Unit V	Drug stability: Reaction kinetics: zero, pseudo-zero, first & second	10 hours
	order, units of basic rate constants, determination of reaction order.	
	Physical and chemical factors influencing the chemical degradation	
	of pharmaceutical product: temperature, solvent, ionic strength,	
	dielectric constant, specific & general acid base catalysis, Simple	
	numerical problems. Stabilization of medicinal agents against	
	common reactions like hydrolysis & oxidation. Accelerated stability	
	testing in expiration dating of pharmaceutical dosage forms.	
	Photolytic degradation and its prevention	

- 1. Physical Pharmacy by Alfred Martin, Sixth edition
- 2. Experimental pharmaceutics by Eugene, Parott.
- 3. Tutorial pharmacy by Cooper & Gunn.
- 4. Stocklosam J. Pharmaceutical calculations, Lea & Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, Marcel Dekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical dosage forms. Disperse systems, volume 1,2,3Marcel Dekkar Inc.
- 7. Physical Pharmaceutics by Ramasamy C, and Manavalan R.

BP407P	PHYSICAL PHARMACEUTICS- II (Practical)	4 Hours/Week	
S.No.	Experiment		
1	Determination of particle size, particle size distribution usin	g sieving method	
2	Determination of particle size, particle size distribution usin	g microscopic method	
3	Determination of bulk density, true density and porosity		
4	Determine the angle of repose and influence of lubricant on	angle of repose	
5	Determination of viscosity of liquid using Ostwald's viscon	neter	
6	Determination sedimentation volume with effect of differen	t suspending agent	
7	Determination sedimentation volume with effect of different concentration of single		
	suspending agent		
8	Determination of viscosity of semisolid by using Brookfield	l viscometer	
9	Determination of reaction rate constant first order		
10	Determination of reaction rate constant second order	Determination of reaction rate constant second order	
11	Accelerated stability studies		

BP404T	PHARMACOLOGY-I (THEORY)	45 Hours	
Scope: The main purpose of the subject is to understand what drugs do to the living organisms and how their effects can be applied to therapeutics. The subject covers the information about the drugs like, mechanism of action, physiological and biochemical effects (pharmacodynamics) as well as absorption, distribution, metabolism and excretion (pharmacokinetics) along with the adverse effects, clinical uses, interactions, doses, contraindications and routes of administration of different classes of drugs.			
	COURSE OUTCOMES		
CO 1	Understand the basic principles of pharmacology and their apstudy of drug actions and interactions in biological systems.	oplication in the	
CO 2	Analyze the pharmacodynamics and pharmacokinetics of drugs absorption, distribution, metabolism, and excretion (ADME).	s, including their	
CO 3	Discuss the mechanisms of drug-receptor interactions, enzymosignal transduction pathways involved in pharmacological response.		
CO 4	Evaluate the pharmacological effects and therapeutic uses of dru autonomic nervous system, central nervous system, and other ta		
CO 5	Apply pharmacological principles in the rational selection and optimization of drug therapy regimens for specific disease conditions.		
	COURSE CONTENT		
Unit I	1. General Pharmacology	08 hours	
	a. Introduction to Pharmacology- Definition, historical landmarks and scope of pharmacology, nature and source of drugs, essential drugs concept and routes of drug administration, Agonists, antagonists (competitive and noncompetitive), spare receptors, addiction, tolerance, dependence, tachyphylaxis, idiosyncrasy, allergy.		
	b. Pharmacokinetics- Membrane transport, absorption, distribution, metabolism and excretion of drugs. Enzyme induction, enzyme inhibition, kinetics of elimination		
Unit II	General Pharmacology	12 hours	
	 c. Pharmacodynamics- Principles and mechanisms of drug action. Receptor theories and classification of receptors, regulation of receptors. drug receptors interactions signal transduction mechanisms, G-protein-coupled receptors, ion channel receptor, transmembrane enzyme linked receptors, transmembrane JAK-STAT binding receptor and receptors that regulate transcription factors, dose response relationship, therapeutic index, combined effects of drugs and factors modifying drug action. d. Adverse drug reactions. 		
	-	-1	

	e. Drug interactions (pharmacokinetic and pharmacodynamic)	
	f. Drug discovery and clinical evaluation of new drugs -Drug	
	discovery phase, preclinical evaluation phase, clinical trial	
	phase, phases of clinical trials and pharmacovigilance.	
Unit III	Pharmacology of drugs acting on peripheral nervous	10 hours
	system	10 hours
	a. Organization and function of ANS.	
	b. Neurohumoral transmission-transmission and classification of neurotransmitters.	
	c. Parasympathomimetics, Parasympatholytics,	
	Sympathomimetics, sympatholytics.	
	d. Neuromuscular blocking agents and skeletal muscle	
	relaxants (peripheral).	
	e. Local anesthetic agents.	
	f. Drugs used in myasthenia gravis and glaucoma	
Unit IV	Pharmacology of drugs acting on central nervous system	08 hours
	a. Neurohumoral transmission in the C.N.S. special emphasis	
	on importance of various	
	b. neurotransmitters like with GABA, Glutamate, Glycine,	
	serotonin, dopamine. b. General anesthetics and pre-	
	anesthetics.	
	c. Sedatives, hypnotics and centrally acting muscle relaxants.	
	d. Anti-epileptics; Alcohols and disulfiram	
Unit V	Pharmacology of drugs acting on central nervous system	07 hours
	a. Psychopharmacological agents: Antipsychotics,	
	antidepressants, anti-anxiety agents, anti-manics and	
	hallucinogens.	
	b. Drugs used in Parkinsons disease and Alzheimer's disease. c.	
	CNS stimulants and nootropics.	
	d. Opioid analgesics and antagonists	
	e. Drug addiction, drug abuse, tolerance and dependence.	
	RECOMMENDED BOOKS (LATEST EDITIONS)	

- 1. Introduction to chemical engineering Walter L Badger & Julius Banchero, Latest
- 2. edition.
- 3. Solid phase extraction, Principles, techniques and applications by Nigel J.K. Simpson-Latest edition.
- 4. Unit operation of chemical engineering Mcabe Smith, Latest edition.
- 5. Pharmaceutical engineering principles and practices C.V.S Subrahmanyam et al.,
- 6. Remington practice of pharmacy- Martin, Latest edition.
- 7. Theory and practice of industrial pharmacy by Lachmann., Latest edition.
- 8. Physical pharmaceutics- C.V.S Subrahmanyam et al., Latest edition.
- 9. Cooper and Gunn's Tutorial pharmacy, S.J. Carter, Latest edition.

BP408P	PHARMACOLOGY-I (Practical)	4 Hours/Week
S.No.	Experiment	
1	Introduction to experimental pharmacology	
2	Commonly used instruments in experimental pharmace	ology
3	Study of common laboratory animals	
4	Maintenance of laboratory animals as per CPCSEA gu	idelines
5	Common laboratory techniques	
6	Study of different routes of drug administration in mice/rats	
7	Effect of hepatic microsomal enzyme inducers on phenobarbitone sleeping time	
8	Effect of drugs on ciliary motility of frog oesophagus	
9	Effect of drugs on rabbit eye	
10	Effects of skeletal muscle relaxants using rota-rod apparatus	
11	Effect of drugs on locomotor activity using actophotometer	
12	Anticonvulsant effect of drugs by MES and PTZ method	
13	Study of stereotype and anti-catatonic activity of drugs	
14	Study of anxiolytic activity of drugs using rats/mice	
15	Study of local anaesthetics by different methods	

Note: All laboratory techniques and animal experiments are demonstrated by simulated experiments by softwares and videos

BP405T	PHARMACOGNOSY AND PHYTOCHEMISTRY I (Theory)	45 Hours	
Scope: The	e subject involves the fundamentals of Pharmacognosy like scope	, classification of	
_	crude drugs, their identification and evaluation, phytochemicals present in them and their		
medicinal p	roperties.		
	COURSE OUTCOMES		
	Acquire comprehensive knowledge of the principles and met	hods used in the	
CO 1	evaluation of crude drugs, including botanical identification	n, phytochemical	
	analysis, and pharmacological testing.	11	
CO 2	Understand the principles and methods of cultivating and oplants, including best agricultural practices and sustain	_	
002	techniques.	able harvesting	
	Comprehend the fundamental principles and techniques	of plant tissue	
CO 3		yogenesis, and	
	organogenesis.		
CO 4	Gain comprehensive knowledge of various alternative syste		
	including their philosophies, principles, and therapeutic appr Evaluate the pharmacological properties and therapeut		
CO 5	medicinal plants and their derived phytoconstituents	de potential of	
	COURSE CONTENT		
Unit I	Introduction to Pharmacognosy:	401	
	(a) Definition, history, scope and development of	10 hours	
	Pharmacognosy		
	(b) Sources of Drugs – Plants, Animals, Marine & Tissue	!	
	culture		
	(c) Organized drugs, unorganized drugs (dried latex, dried juices, dried extracts, gums and		
	mucilages, oleoresins and oleo- gum -resins).		
	Classification of drugs:		
	Alphabetical, morphological, taxonomical, chemical	,	
	pharmacological, chemo and sero		
	taxonomical classification of drugs Quality control of Drugs of Natural Origin:		
	Adulteration of drugs of natural origin. Evaluation by	,	
	organoleptic, microscopic, physical,		
	chemical and biological methods and properties. Quantitative		
	microscopy of crude drugs including lycopodium spore	!	
	method, leafconstants, camera lucida and diagrams of		
	microscopic objects to scale with camera lucida.		
Unit II	Cultivation, Collection, Processing and storage of drugs of	1 () () ()	
	natural origin: Cultivation and Collection of drugs of natural		
	origin Factors influencing cultivation of medicinal plants. Plant hormones and their applications. Polyploidy, mutation		
	and hybridization with reference to medicinal plants		
	Conservation of medicinal plants		
Unit III	Plant tissue culture: Historical development of plant tissue	07 hours	
		o/ Hours	

	culture, types of cultures, Nutritional requirements, growth		
	and their maintenance. Applications of plant tissue culture in		
	pharmacognosy. Edible vaccines		
Unit IV	Pharmacognosy in various systems of medicine:	10 hours	
	Role of Pharmacognosy in allopathy and traditional systems of		
	medicine namely, Ayurveda, Unani, Siddha, Homeopathy and		
	Chinese systems of medicine.		
	Introduction to secondary metabolites: Definition,		
	classification, properties and test for identification of		
	Alkaloids, Glycosides, Flavonoids, Tannins, Volatile oil and		
	Resins		
Unit V	Study of biological source, chemical nature and uses of drugs	08 hours	
	of natural origin containing following drugs		
	Plant Products: Fibers - Cotton, Jute, Hemp Hallucinogens,		
	Teratogens, Natural allergens		
	Primary metabolites: General introduction, detailed study with		
	respect to chemistry, sources, preparation, evaluation,		
	preservation, storage, therapeutic used and commercial utility as		
	Pharmaceutical Aids and/or Medicines for the following Primary		
	metabolites:		
	Carbohydrates: Acacia, Agar, Tragacanth, Honey Proteins and		
	Enzymes: Gelatin, casein, proteolytic enzymes (Papain,		
	bromelain, serratiopeptidase, urokinase, streptokinase, pepsin).		
	Lipids (Waxes, fats, fixed oils): Castor oil, Chaulmoogra oil,		
	Wool Fat, Bees Wax Marine Drugs: Novel medicinal agents from		
	marine sources		
	RECOMMENDED BOOKS (LATEST EDITIONS)		

- 1. W.C. Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Tyler, V.E., Brady, L.R. and Robbers, J.E., Pharmacognosy, 9th Edn., Lea and Febiger, Philadelphia, 1988.
- 3. Text Book of Pharmacognosy by T.E. Wallis
- 4. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 5. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 6. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 7. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi. 2007
- 8. Practical Pharmacognosy: C.K. Kokate, Purohit, Gokhlae
- 9. Anatomy of Crude Drugs by M.A. IyengarCooper and Gunn's Tutorial pharmacy, S.J. Carter, Latest edition.

BP409P	Pharmacognosy and Phytochemistry I – (Practical)	4 Hours/Week	
S.No.	Experiment		
1	Analysis of crude drugs by chemical tests		
2	Determination of stomatal number and index	Determination of stomatal number and index	
3	Determination of vein islet number, vein islet termination and palisade ratio		
4	Determination of size of starch grains, calcium oxalate crystals		
5	Determination of Fiber length and width		
6	Determination of number of starch grains by Lycopodium spore method		
7	Determination of Ash value		
8	Determination of Extractive values of crude drugs		
9	Determination of moisture content of crude drugs		
10	Determination of swelling index and foaming		

BP501T	MEDICINAL CHEMISTRY - II (Theory)	45 Hours	
	bject is designed to impart fundamental knowledge on the structure		
	therapeutic value of drugs. The subject emphasizes on structure activity relationships of drugs,		
	physicochemical properties and metabolism of drugs. The syllabus a	ilso emphasizes	
on chemical sy	nthesis of important drugs under each class. COURSE OUTCOMES		
00.4			
CO 1	Analyze the chemical structures of various classes of drugs.		
CO 2	Understand the mechanisms of action and SAR of medicinal co		
CO 3	Evaluate the pharmacokinetic properties and metabolism of d	rugs.	
CO 4	Explain the synthesis and design of new drug molecules.		
CO 5	Apply knowledge of medicinal chemistry in drug discovery pr	ocesses.	
	COURSE CONTENT		
Study of the	development of the following classes of drugs, Classification,	mechanism of	
	of drugs mentioned in the course, Structure activity relationsh		
	as specified in the course and synthesis of drugs superscripted		
Unit I	Antihistaminic agents: Histamine, receptors and their distribution		
	in the human body	10 Hours	
	H1-antagonists: Diphenhydramine hydrochloride ³		
	Dimenhydrinate, Doxylamines cuccinate, Clemastine fumarate		
	Diphenylphyraline hydrochloride, Tripelenamine hydrochloride		
	Chlorcyclizine hydrochloride, Meclizine hydrochloride, Buclizin hydrochloride, Chlorpheniramine maleate, Triprolidin		
	hydrochloride*, Phenidamine tartarate, Promethazine hydrochloride*, Trimeprazine tartrate, Cyproheptadine		
	hydrochloride, Azatidine maleate, Astemizole, Loratadine, Cetirizine,		
	Levocetrazine Cromolyn sodium		
	H2-antagonists: Cimetidine*, Famotidine, Ranitidin.		
	Gastric Proton pump inhibitors: Omeprazole, Lansoprazole,		
	Rabeprazole, Pantoprazole		
	Anti-neoplastic agents:		
	Alkylating agents: Meclorethamine*, Cyclophosphamid	e	
	Melphalan, Chlorambucil, Busulfan, Thiotepa Antimetabolites: Mercaptopurine*, Thioguanine, Fluorouraci	1	
	Floxuridine, Cytarabine, Methotrexate*, Azathioprine	·,	
	Antibiotics: Dactinomycin, Daunorubicin, Doxorubicin, Bleomycin	$_{\rm n}$	
	Plant products: Etoposide, Vinblastin sulphate, Vincristin sulphat		
	Miscellaneous: Cisplatin, Mitotane.		
Unit II	Anti-anginal: Vasodilators: Amyl nitrite, Nitroglycerin*	. 10 hours	
	Pentaerythritol tetranitrate, Isosorbide dinitrite [*]	, To nours	
	Dipyridamole.		
	Calcium channel blockers: Verapamil, Bepridil		
	hydrochloride, Diltiazem hydrochloride, Nifedipine, Amlodipine	2,	
	Felodipine, Nicardipine, Nimodipine.	k	
	Diuretics: Carbonic anhydrase inhibitors: Acetazolamide	,	
	Methazolamide, Dichlorphenamide. Thiazides: Chlorthiazide*, Hydrochlorothiazide	<u>, </u>	
	Hydroflumethiazide, Cyclothiazide,	ē,	
	Loop diuretics: Furosemide*, Bumetanide, Ethacrynic acid	ı.	
	Potassium sparing Diuretics: Spironolactone, Triamterene		
	Transcorer, Transcorer,	7 1	

	Amiloride. Osmotic Diuretics: Mannitol		
	Anti-hypertensive Agents: Timolol, Captopril, Lisinopril, Enalapril,		
	Benazepril hydrochloride, Quinapril hydrochloride, Methyldopate		
	hydrochloride, * Clonidine hydrochloride, Guanethidine		
	monosulphate, Guanabenz acetate, Sodium nitroprusside, Diazoxide,		
	Minoxidil, Reserpine, Hydralazine hydrochloride		
II!A III			
Unit III	Anti-arrhythmic Drugs: Quinidine sulphate, Procainamide hydrochloride, Disopyramide phosphate*, Phenytoin sodium,	10 hours	
	Lidocaine hydrochloride, Tocainide hydrochloride, Mexiletine hydrochloride, Lorcainide hydrochloride, Amiodarone, Sotalol.		
	Anti-hyperlipidemic agents: Clofibrate, Lovastatin,		
	Cholesteramine and Cholestipol		
	Coagulant & Anticoagulants: Menadione, Acetomenadione,		
	Warfarin*, Anisindione, clopidogrel		
	Congestive Heart Failure: Digoxin, Digitoxin, Nesiritide, Bosentan,		
	Tezosentan.		
Unit IV	Drugs acting on Endocrine system: Nomenclature,	08 hours	
	Stereochemistry and metabolism of steroids		
	Sex hormones: Testosterone, Nandralone, Progestrones, Oestriol,		
	Oestradiol, Oestrione, Diethyl stilbestrol.		
	Drugs for erectile dysfunction: Sildenafil, Tadalafil.		
	Oral contraceptives: Mifepristone, Norgestril, Levonorgestrol		
	Corticosteroids: Cortisone, Hydrocortisone, Prednisolone,		
	Betamethasone, Dexamethasone		
	Thyroid and antithyroid drugs: L-Thyroxine, L-Thyronine,		
	Propylthiouracil, Methimazole.		
Unit V	Antidiabetic agents: Insulin and its preparations	07 hours	
	Sulfonyl ureas: Tolbutamide*, Chlorpropamide, Glipizide,		
	Glimepiride. Biguanides: Metformin.		
	Thiazolidinediones: Pioglitazone, Rosiglitazone.		
	Meglitinides: Repaglinide, Nateglinide.		
	Glucosidase inhibitors: Acrabose, Voglibose.		
	Local Anesthetics: SAR of Local anesthetics		
	Benzoic Acid derivatives; Cocaine, Hexylcaine, Meprylcaine,		
	Cyclomethycaine, Piperocaine.		
	Amino Benzoic acid derivatives: Benzocaine*, Butamben,		
	Procaine*, Butacaine, Propoxycaine, Tetracaine, Benoxinate.		
	Lidocaine/Anilide derivatives: Lignocaine, Mepivacaine,		
	Prilocaine, Etidocaine.		
	Miscellaneous: Phenacaine, Diperodon, Dibucaine. *		
	RECOMMENDED BOOKS (LATEST EDITIONS)		

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1to 5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I. Vogel.

BP 502 T	Industrial Pharmacy I (Theory)	45 Hours		
Scope: Cou	Scope : Course enables the student to understand and appreciate the influence of			
*	pharmaceutical additives and various pharmaceutical dosage forms on the			
performan	ce of the drug product.			
	COURSE OUTCOMES			
CO 1	Understand and apply preformulation principles to evaluate dru	ig properties for		
	predicting formulation challenges and ensuring stability.	11		
CO 2	Formulate, manufacture, and perform quality control on tablets	and liquid orals,		
	understanding their production processes and challenges.			
CO 3	Understand production processes and quality control for hard capsules and the formulation and manufacturing of pellets.	and sort gelatin		
	Gain knowledge in the formulation, production, and quality cont	rol of parenteral		
CO 4	and ophthalmic products, focusing on aseptic processing and st	-		
	Formulate and prepare cosmetic products and pharmaceutic	•		
CO 5	understand packaging materials science and quality control for p			
	and safety.			
	COURSE CONTENT			
Unit I	Preformulation Studies: Introduction to preformulation, goals and	d 07 hours		
	objectives, study of physicochemical characteristics of drug	g 07 Hours		
	substances.			
	Physical properties: Physical form (crystal & amorphous), particle size, shape, flow properties, solubility profile (pKa, pH, partition			
	coefficient), polymorphism			
	Chemical Properties: Hydrolysis, oxidation, reduction, racemization,			
	polymerization BCS classification of drugs & its significant			
	Application of preformulation considerations in the development of			
	solid, liquid oral and parenteral dosage forms and its impact on stability of dosage forms.			
Unit II	Tablets: Introduction, ideal characteristics of tablets, classification of	of 10 h		
	tablets. Excipients, Formulation of tablets, granulation methods	$\frac{1}{5}$, 10 hours		
	compression and processing problems. Equipments and tablet tooling			
	Tablet coating: Types of coating, coating materials, formulation of coating composition, methods of coating, equipment employed and			
	defects in coating.			
	Quality control tests: In process and finished product tests			
	Liquid orals: Formulation and manufacturing consideration of syrup			
	and elixirs suspensions and emulsions; Filling and packaging	5;		
Unit III	evaluation of liquid orals official in pharmacopoeia Capsules:	001		
	Hard gelatin capsules: Introduction, Production of hard gelating	n 08 hours		
	capsule shells. Size of capsules, Filling, finishing and special technique	S		
	of formulation of hard gelatin capsules, manufacturing defects. In	n		
	process and final product quality control tests for capsules. Soft gelatin capsules: Nature of shell and capsule content, size of	.f		
	capsules, importance of base adsorption and minim/gram factors			
	production, in process and final product quality control tests. Packing			
	storage and stability testing of soft gelatin capsules and thei			
	applications.			

	Pellets: Introduction, formulation requirements, pelletization		
	process, Equipments for manufacture of pellets		
Unit IV	Parenteral Products: 10 hours		
	Definition, types, advantages and limitations. Preformulation factors		
	and essential requirements, vehicles, additives, importance of		
	isotonicity Production procedure, production facilities and controls,		
	aseptic processing. Formulation of injections, sterile powders, large		
	volume parenteral and lyophilized products. Containers and closures		
	selection, filling and sealing of ampoules, vials and infusion fluids.		
	Quality control tests of parenteral products.		
	Ophthalmic Preparations:		
	Introduction, formulation considerations; formulation of eye drops,		
	eye ointments and eye lotions; methods of preparation; labeling,		
	containers; evaluation of ophthalmic preparations		
Unit V	Cosmetics:	10 hours	
	Formulation and preparation of the following cosmetic preparations:		
	lipsticks, shampoos, cold cream and vanishing cream, tooth pastes,		
	hair dyes and sunscreens.		
	Pharmaceutical Aerosols:		
	Definition, propellants, containers, valves, types of aerosol systems;		
	formulation and manufacture of aerosols; Evaluation of aerosols;		
	Quality control and stability studies.		
	Packaging Materials Science:		
	Materials used for packaging of pharmaceutical products, factors		
	influencing choice of containers, legal and official requirements for		
	containers, stability aspects of packaging materials, quality control		
	tests.		

- 1. Pharmaceutical dosage forms Tablets, volume 1 -3 by H.A. Liberman, Leon Lachman&J.B. Schwartz
- 2. Pharmaceutical dosage form Parenteral medication vol- 1&2 by Liberman &Lachman
- $3. \ \ Pharmaceutical\ dosage\ form\ disperse\ system\ VOL-1\ by\ Liberman\ \&\ Lachman$
- 4. Modern Pharmaceutics by Gilbert S. Banker & C.T. Rhodes, 3rd Edition
- 5. Remington: The Science and Practice of Pharmacy, 20th edition PharmaceuticalScience (RPS)
- 6. Theory and Practice of Industrial Pharmacy by Liberman & Lachman
- 7. Pharmaceutics- The science of dosage form design by M.E. Aulton, Churchilllivingstone, Latest edition
- 8. Introduction to Pharmaceutical Dosage Forms by H. C. Ansel, Lea &Febiger, Philadelphia, 5thedition, 2005
- 9. Drug stability Principles and practice by Cartensen & C.J. Rhodes, 3rd Edition, Marcel Dekker Series, Vol 107.

BP502P	Industrial Pharmacy I (Practical)	4 Hours/Week
S.No.	Experiment	
1	Preformulation studies on Paracetamol/Aspirin/or any o	other drug
2	Preparation and evaluation of Paracetamol tablets	
3	Preparation and evaluation of Aspirin tablets	
4	Coating of tablets- film coating of tables/granules	
5	Preparation and evaluation of Tetracycline capsules	
6	Preparation of Calcium Gluconate injection	
7	Preparation of Ascorbic Acid injection	
8	Quality control test of (as per IP) marketed tablets and o	capsules
9	Preparation of Eye drops and Eye ointments	
10	Preparation of Creams (Cold / Vanishing cream)	
11	Evaluation of Glass containers (as per IP)	

	DHADMACOLOCV II (Theowy)	45 Hours	
BP503T	PHARMACOLOGY-II (Theory)	15 110415	
_	Scope: This subject is intended to impart the fundamental knowledge on various		
	aspects (classification, mechanism of action, therapeutic effects, clinical uses, side		
	contraindications) of drugs acting on different systems of body	y and in	
addition, er	mphasis on the basic concepts of bioassay. COURSE OUTCOMES		
CO 1		rigal greateme	
	Understand the mechanisms of drug action on various physiolog	<u> </u>	
CO 2	Analyze the pharmacological effects and therapeutic uses of drug		
CO 3	Evaluate the adverse effects and toxicity profiles of pharmacolog		
CO 4	Explain the principles of bioassay and its application in assaying	varying	
	drugs.		
CO 5	Apply pharmacological concepts in clinical practice and drug the	erapy.	
	COURSE CONTENT		
Unit I	1. Pharmacology of drugs acting on cardio vascular system	08 hours	
	a. Introduction to hemodynamic & electrophysiology of heart.	U8 nours	
	b. Drugs used in congestive heart failure		
	c. Anti-hypertensive drugs.		
	d. Anti-anginal drugs.		
	e. Anti-arrhythmic drugs.		
** ** **	f. f. Anti-hyperlipidemic drugs.		
Unit II	1. Pharmacology of drugs acting on cardio vascular system	12 hours	
	a. Drug used in the therapy of shock.b. Hematinics, coagulants and anticoagulants.		
	c. Fibrinolytics and anti-platelet drugs		
	d. Plasma volume expanders		
	2. Pharmacology of drugs acting on urinary system		
	a. Diuretics		
	b. Anti-diuretics.		
Unit III	Autocoids and related drugs	10 hours	
	a. Introduction to autacoids and classification		
	b. Histamine, 5-HT and their antagonists.		
	c. Prostaglandins, Thromboxanes and Leukotrienes.d. Angiotensin, Bradykinin and Substance P.		
	d. Angiotensin, Bradykinin and Substance P. e. Non-steroidal anti-inflammatory agents		
	f. Anti-gout drugs		
	a. Antirheumatic drugs		
Unit IV	Pharmacology of drugs acting on endocrine system	08 hours	
	a. Basic concepts in endocrine pharmacology.		
	b. Anterior Pituitary hormones- analogues and their inhibitors.		
	c. Thyroid hormones- analogues and their inhibitors.		
	d. Hormones regulating plasma calcium level- Parathormone,		
	Calcitonin and Vitamin-D.		
	e. Insulin, Oral Hypoglycemic agents and glucagon.		
	f. ACTH and corticosteroids.		

Unit V	Pharmacology of drugs acting on endocrine system	07 hours
	a. Androgens and Anabolic steroids.	
	b. Estrogens, progesterone and oral contraceptives.	
	c. Drugs acting on the uterus.	
	6. Bioassay	
	a. Principles and applications of bioassay.	
	b. Types of bioassay	
	c. Bioassay of insulin, oxytocin, vasopressin, ACTH, d-	
	tubocurarine, digitalis, histamine and 5-HT	
DECOMMENDED DOOMS (LAMECE EDIMIONS)		

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill.
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A.
- 5. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs, The Point
- 6. Lippincott Williams & Wilkins.
- 7. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews- Pharmacology.
- 8. K.D. Tripathi. Essentials of Medical Pharmacology, JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 9. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher
- 10. Modern Pharmacology with clinical Applications, by Charles R. Craig& Robert.
- 11. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.
- 12. Kulkarni SK. Handbook of experimental pharmacology. Vallabh Prakashan.

BP507P	PHARMACOLOGY-I (Practical)	4 Hours/Week
S.No.	Experiment	
1	Introduction to in-vitro pharmacology and physiological	l salt solutions
2	Effect of drugs on isolated frog heart	
3	Effect of drugs on blood pressure and heart rate of dog	
4	Study of diuretic activity of drugs using rats/mice	
5	DRC of acetylcholine using frog rectus abdominis muscl	e
6	Effect of physostigmine and atropine on DRC of acetylcholine	
7	Bioassay of histamine using guinea pig ileum by matching method	
8	Bioassay of oxytocin using rat uterine horn by interpolation method	
9	Bioassay of serotonin using rat fundus strip by three-point bioassay	
10	Bioassay of acetylcholine using rat ileum/colon by four-point bioassay	
11	Determination of PA2 value of prazosin using rat anococcygeus muscle (by Schilds plot	
	method)	
12	Determination of PD2 value using guinea pig ileum	
13	Effect of spasmogens and spasmolytics using rabbit jejunum	
14	Anti-inflammatory activity of drugs using carrageenan induced paw-edema model	
15	Analgesic activity of drug using central and peripheral methods	

Note: All laboratory techniques and animal experiments are demonstrated by simulated experiments by softwares and videos

BP504T	PHARMACOGNOSY AND PHYTOCHEMISTRY II (Theory)	45 Hours		
Scope: The	e, classification			
of crude dr	of crude drugs, their identification and evaluation, phytochemicals present in them and			
their medicinal properties.				
COURSE OUTCOMES				
CO 1	Gain in-depth knowledge of the metabolic pathways involved is	•		
00 1	including photosynthesis, respiration, and secondary metaboli	•		
CO 2	Understand the therapeutic applications of secondary metabotheir roles in traditional and modern medicine. Learn pharmacological properties and Commercial applications.			
20.0	Acquire a knowledge of phytoconstituents, their types, and	their roles in		
CO 3	plants, including alkaloids, flavonoids, terpenoids, glycosides, a			
CO 4	Apply techniques for the extraction, isolation, and characterizat compounds.	cion of bioactive		
CO 5	Gain a solid understanding of the basic principles of phytochem			
	the study of bioactive compounds in plants and their chemical COURSE CONTENT	properties.		
Unit I Metabolic pathways in higher plants and their 07 hours				
Uniti	determination	07 hours		
	a) Brief study of basic metabolic pathways and formation of			
	different secondary metabolites			
	through these pathways- Shikimic acid pathway, Acetate			
	pathways and Amino acid pathway.			
	b) Study of utilization of radioactive isotopes in the			
	investigation of Biogenetic studies			
Unit II	General introduction, composition, chemistry & chemical	14 hours		
	classes, biosources, therapeutic	14 nours		
	uses and commercial applications of following			
	secondary metabolites:			
	Alkaloids: Vinca, Rauwolfia, Belladonna, Opium, Phenylpropanoids and Flavonoids: Lignans, Tea, Ruta			
	Steroids, Cardiac Glycosides & Triterpenoids: Liquorice,			
	Dioscorea, Digitalis			
	Volatile oils: Mentha, Clove, Cinnamon, Fennel, Coriander,			
	Tannins: Catechu, Pterocarpus			
	Resins: Benzoin, Guggul, Ginger, Asafoetida, Myrrh,			
	Colophony Changidan Sonna Alaca Bittar Almand			
	Glycosides: Senna, Aloes, Bitter Almond Iridoids, Other terpenoids & Naphthaquinones: Gentian,			
	Artemisia, taxus, carotenoids			
Unit III	Isolation, Identification and Analysis of	06 h a		
	Phytoconstituents	06 hours		
	a) Terpenoids: Menthol, Citral, Artemisin			
	b) Glycosides: Glycyrhetinic acid & Rutin			
	c) Alkaloids: Atropine, Quinine, Reserpine, Caffeine			

	d) Resins: Podophyllotoxin, Curcumin	
Unit IV	Industrial production, estimation and utilization of the	10 hours
	following phytoconstituents: Forskolin, Sennoside,	
	Artemisinin, Diosgenin, Digoxin, Atropine, Podophyllotoxin,	
	Caffeine, Taxol, Vincristine and Vinblastine	
II!+ V		00.1
Unit V	Basics of Phytochemistry Modern methods of extraction,	08 hours
Unit	application of latest techniques like Spectroscopy,	08 nours
Unit v		08 nours
Unit v	application of latest techniques like Spectroscopy,	08 nours

- 1. W.C. Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 3. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 4. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 5. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007
- 6. Herbal Cosmetics by H. Pande, Asia Pacific Business press, Inc, New Delhi.
- 7. A.N. Kalia, Textbook of Industrial Pharmacognosy, CBS Publishers, New Delhi, 2005.
- 8. R Endress, Plant cell Biotechnology, Springer-Verlag, Berlin, 1994.
- 9. Pharmacognosy & Pharmacobiotechnology. James Bobbers, Marilyn KS, VE Tylor.
- 10. The formulation and preparation of cosmetic, fragrances and flavours. 11. Remington's Pharmaceutical sciences. 12. Text Book of Biotechnology by Vyas and Dixit.
- 11. Text Book of Biotechnology by R.C. Dubey

BP508P	Pharmacognosy and Phytochemistry II - (Practical)	4 Hours/Week
S.No.	Experiment	
1	Morphology, histology, and powder characteristics & extraction & Cinchona, Cinnamon, Senna, Clove, Ephedra, Fennel, and Coriander	
2	Exercise involving isolation & detection of active principles: Caffeine from tea dust Diosgenin from Dioscorea Atropine from Belladonna Sennosides from Senna	
3	Separation of sugars by Paper chromatography	
4	TLC of herbal extract	
5	Distillation of volatile oils and detection of phytoconstituents by T	LC
6	Analysis of crude drugs by chemical tests: Asafoetida, Benzoin, Col Myrrh	ophony, Aloes,

B. PHARM (SEMESTER - VI)

BP601T MEDICINAL CHEMISTRY-III (Theory) 45 Hours

Scope: This subject is designed to impart fundamental knowledge on the structure, chemistry and therapeutic value of drugs. The subject emphasis on modern techniques of rational drug design like quantitative structure-activity relationship (QSAR), Prodrug concept, combinatorial chemistry and Computer-aided drug design (CADD). The subject also emphasizes on the chemistry, mechanism of action, metabolism, adverse effects, Structure Activity Relationships (SAR), therapeutic uses and synthesis of important drugs.

COURSE OUTCOMES Analyze advanced chemical structures of medicinal compounds.

CO 2	Understand drug design principles for specific therapeutic targets.	
CO 3	Evaluate the chemical classes and mechanism of action of drugs.	
CO 4	Explain the structure-activity relationships (SAR) in drug discovery.	

CO 5 Apply computational methods in medicinal chemistry research.

CO 1

COURSE CONTENT

Study of the development of the following classes of drugs, Classification, mechanism of action, uses of drugs mentioned in the course, Structure-activity relationship of selective class of drugs as specified in the course and synthesis of drugs superscripted by (*)

Unit I	Antibiotics	10
	Historical background, Nomenclature, Stereochemistry,	hours
	Structure-activity relationship, Chemical degradation	Hours
	classification and important products of the following classes.	
	β-Lactam antibiotics: Penicillin, Cephalosporins, β- Lactamase	
	inhibitors, Monobactams	
	Aminoglycosides: Streptomycin, Neomycin, Kanamycin	
	Tetracyclines: Tetracycline, Oxytetracycline, Chlortetracycline,	
	Minocycline, Doxycycline	
Unit II	Antibiotics	10
	Historical background, Nomenclature, Stereochemistry, Structure-	10
	activity relationship, Chemical degradation classification and	hours
	important products of the following classes.	
	Macrolide: Erythromycin Clarithromycin, Azithromycin.	
	Miscellaneous: Chloramphenicol*, Clindamycin.	
	Prodrugs : Basic concepts and application of prodrugs design.	

	Antimalarials: Etiology of malaria.	
	Quinolines : SAR, Quinine sulfate, Chloroquine*,	
	Amodiaquine, Primaquine phosphate, Pamaquine*, Quinacrine	
	hydrochloride, Mefloquine.	
	Biguanides and dihydro triazines: Cycloguanil pamoate,	
	Proguanil.	
	Miscellaneous : Pyrimethamine, Artesunate, Artemether,	
	Atovaquone.	
Unit III	Anti-tubercular Agents	4.0
	Synthetic anti-tubercular agents: Isoniazid*, Ethionamide,	10
	Ethambutol, Pyrazinamide, Para amino salicylic acid. *	hours
	Anti-tubercular antibiotics: Rifampicin, Rifabutin, Cycloserine	
	Streptomycin, Capreomycin sulfate.	
	Urinary tract anti-infective agents	
	Quinolones: SAR of quinolones, Nalidixic Acid, Norfloxacin,	
	Enoxacin, Ciprofloxacin*, Ofloxacin, Lomefloxacin, Sparfloxacin,	
	Gatifloxacin, Moxifloxacin	
	Miscellaneous: Furazolidone, Nitrofurantoin*, Methenamine.	
	Antiviral agents:	
	Amantadine hydrochloride, Rimantadine hydrochloride,	
	Idoxuridine trifluoride, Acyclovir*, Ganciclovir, Zidovudine,	
	Didanosine, Zalcitabine, Lamivudine, Loviride, Delavirdine,	
	Ribavirin, Saquinavir, Indinavir, Ritonavir.	
Unit IV	Antifungal agents:	08
	Antifungal antibiotics: Amphotericin-B, Nystatin, Natamycin,	hours
	Griseofulvin.	nours
	Synthetic Antifungal agents: Clotrimazole, Econazole,	
	Butoconazole, Oxiconazole Tioconazole, Miconazole*,	
	Ketoconazole, Terconazole, Itraconazole, Fluconazole, Naftifine	
	hydrochloride, Tolnaftate*.	
	Anti-protozoal Agents: Metronidazole*, Tinidazole, Ornidazole,	
	Diloxanide, Iodoquinol, Pentamidine Isethionate, Atovaquone,	
	Eflornithine.	
	Eflornithine. Anthelmintics: Diethylcarbamazine citrate*. Thiabendazole.	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole,	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole,	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine,	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin.	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin. Sulphonamides and Sulfones Historical development, chemistry, classification and SAR of	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin. Sulphonamides and Sulfones	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin. Sulphonamides and Sulfones Historical development, chemistry, classification and SAR of Sulfonamides: Sulphamethizole, Sulfisoxazole, Sulphamethizine,	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin. Sulphonamides and Sulfones Historical development, chemistry, classification and SAR of Sulfonamides: Sulphamethizole, Sulfisoxazole, Sulphamethizine, Sulfacetamide*, Sulphapyridine, Sulfamethoxazole*, Sulphadiazine,	
	Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantel, Ivermectin. Sulphonamides and Sulfones Historical development, chemistry, classification and SAR of Sulfonamides: Sulphamethizole, Sulfisoxazole, Sulphamethizine, Sulfacetamide*, Sulphapyridine, Sulfamethoxazole*, Sulphadiazine, Mafenide acetate, Sulfasalazine.	

Unit V	Introduction to Drug Design	07
	Various approaches used in drug design.	hours
	Physicochemical parameters used in quantitative structure activity	
	relationship (QSAR) such as partition coefficient, Hammet's	
	electronic parameter, Tafts steric parameter and Hansch analysis.	
	Pharmacophore modelling and docking techniques.	
	Combinatorial Chemistry: Concept and applications of	
	combinatorial chemistry: solid phase and solution phase synthesis.	

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I.Vogel.

BP607P	MEDICINAL CHEMISTRY-III (Practical)	4 Hours/Week

Scope: Practical physiology is complimentary to the theoretical discussions in physiology. Practicals allow the verification of physiological processes discussed in theory classes through experiments on living tissue, intact animals or normal human beings. This is helpful for developing an insight on the subject.

S.No.	Topic	
I.	Preparation of drugs and intermediates	
1	Sulphanilamide	
2	7-Hydroxy, 4-methyl coumarin	
3	Chlorobutanol	
4	Triphenyl imidazole	
5	Tolbutamide	
6	Hexamine	
II.	Assay of drugs	
1	Isonicotinic acid hydrazide	
2	Chloroquine	
3	Metronidazole	
4	Dapsone	
5	Chlorpheniramine maleate	
6	Benzylpenicillin	
III.	Preparation of medicinally important compounds or intermediates	
	microwave irradiation technique	
IV.	Drawing structures and reactions using chem draw®	
V.	Determination of physicochemical properties such as logP, clogP, MR,	
	Molecular weight, Hydrogen bond donors and acceptors for class of drugs	
	course content using drug design software Drug likeliness screening	
	(Lipinskies RO5)	

BP602T	PHARMACOLOGY-III (Theory)	45 Hours
Scope: This subject is intended to impart the fundamental knowledge on various aspects (classification, mechanism of action, therapeutic effects, clinical uses, side effects and contraindications) of drugs acting on the respiratory and gastrointestinal system, infectious diseases, immuno-pharmacology and in addition, emphasis on the principles of toxicology and chronopharmacology.		
	COURSE OUTCOMES	
CO 1	Analyze the mechanisms of action of advanced pharmacological	agents.
CO 2	Understand the pharmacodynamics and pharmacokinetics of dr	ugs.
CO 3	Evaluate the therapeutic uses and adverse effects of pharmacolo	gical agents.
CO 4	Explain the principles of chronopharmacology and immunophar	macology
CO 5	Apply pharmacological concepts in the treatment of complex dis	seases.
	COURSE CONTENT	
Unit I	1. Pharmacology of drugs acting on Respiratory system	10 hours
	a. Anti -asthmatic drugs	To hours
	b. Drugs used in the management of COPD	
	c. Expectorants and antitussives	
	d. Nasal decongestants	
	e. Respiratory stimulants	
	2. Pharmacology of drugs acting on the Gastrointestinal	
	Tract	
	a. Antiulcer agents.	
	b. Drugs for constipation and diarrhoea.	
	c. Appetite stimulants and suppressants.	
	d. Digestants and carminatives.	
	e. Emetics and anti-emetics.	
Unit II	3. Chemotherapy	10 hours
	a. General principles of chemotherapy.	
	b. Sulfonamides and cotrimoxazole.	
	c.Antibiotics- Penicillin's, cephalosporins, chloramphenicol	
	macrolides, quinolones and fluoroquinolones, tetracycline and	L
Unit III	aminoglycosides	
Unitin	3. Chemotherapy	10 hours
	a. Antitubercular agents b. Antileprotic agents	
	c. Antifungal agents	
	d Antirinal drugs	

d. Antiviral drugs e. Anthelmintics

	f. Antimalarial drugs	
	g. Antiamoebic agents	
Unit IV	3. Chemotherapy	08 hours
	Urinary tract infections and sexually transmitted diseases.	
	Chemotherapy of malignancy.	
	4. Immunopharmacology	
	a. Immunostimulants	
	b. Immunosuppressant	
	Protein drugs, monoclonal antibodies, target drugs to antigen,	
	biosimilars	
Unit V	5. Principles of toxicology	07
	a. Definition and basic knowledge of acute, subacute and chronic	hours
	toxicity.	
	b. Definition and basic knowledge of genotoxicity,	
	carcinogenicity, teratogenicity and mutagenicity	
	c. General principles of treatment of poisoning	
	d. Clinical symptoms and management of barbiturates, morphine,	
	organophosphorus compound and lead, mercury and	
	arsenic poisoning.	
	6. Chronopharmacology	
	a. Definition of rhythm and cycles.	
	b. Biological clock and their significance leading to	
	chronotherapy.	

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs. The Point Lippincott Williams & Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology
- 6. K.D. Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher Modern Pharmacology with clinical Applications, by Charles R. Craig& Robert,
- 8. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata,
- 9. Kulkarni SK. Handbook of experimental pharmacology. VallabhPrakashan,
- 10. N. Udupa and P.D. Gupta, Concepts in Chronopharmacology.

BP608P	PHARMACOLOGY-III (Practical)	4 Hours/Week	
1	Dose calculation in pharmacological experiments.		
2	Antiallergic activity by mast cell stabilization assay.		
3	Study of anti-ulcer activity of a drug using pylorus ligand (SHAY) rat model and		
	NSAIDS-induced ulcer model.		
4	Study of effect of drugs on gastrointestinal motility.		
5	Effect of agonists and antagonists on guinea pig ileum.		
6	Estimation of serum biochemical parameters by using semi- autoanalyzer.		
7	Effect of saline purgative on frog intestine.		
8	Insulin hypoglycaemic effect in rabbit.		
9	Test for pyrogens (rabbit method)		
10	Determination of acute oral toxicity (LD50) of a drug from a given	ren data.	
11	Determination of acute skin irritation / corrosion of a test subs	tance.	
12	Determination of acute eye irritation / corrosion of a test subst	ance	
13	Calculation of pharmacokinetic parameters from a given data		
14	Biostatistics methods in experimental pharmacology (student's	t test, ANOVA)	
15	Biostatistics methods in experimental pharmacology (Chi square test, Wilcoxon		
	Signed Rank test)		
*Experim	*Experiments are demonstrated by simulated experiments/videos		

BP603T	HERBAL DRUG TECHNOLOGY (Theory)	45 Hours		
	Scope: This subject gives the student the knowledge of basic understanding of herbal drug			
	ne quality of raw material, guidelines for quality of herbal drugs, he eeteners, nutraceutical etc. The subject also emphasizes on Good			
	GMP), patenting and regulatory issues of herbal drugs	Manufacturing		
Tractices (j, patenting and regulatory issues of herbar arags			
	COURSE OUTCOMES			
	Gain knowledge of the Herbal medicine, principles and practices			
CO 1	biodynamic agriculture, emphasizing holistic and sustainable far	rming		
	methods.	1.1 1.1		
CO 2	Learn about the Nutraceuticals including their types, sources, an			
CO 2	benefits and role of nutraceuticals in the prevention and manage various diseases	ement of		
	Develop skills in formulating herbal cosmetic products such as c	reams lotions		
CO 3	shampoos ensuring efficacy and stability.	realis, recions,		
CO 4	Explain the regulatory requirements, safety considerations and	Stability		
CO 4	testing of herbal drug drugs			
CO 5	Gain comprehensive knowledge of Schedule T requirements and			
00 5	Manufacturing Practices (GMP) specific to the Indian systems of	medicine		
IIia I	COURSE CONTENT			
Unit I	Herbs as raw materials Definition of herb, herbal medicine, herbal medicinal product,	11 hours		
	herbal drug preparation, Source of Herbs Selection,			
	identification and authentication of herbal materials			
	Processing of herbal raw material.			
	Biodynamic Agriculture			
	Good agricultural practices in cultivation of medicinal plants			
	including Organic farming. Pest and Pest management in			
	medicinal plants: Biopesticides/Bioinsecticides.			
	Indian Systems of Medicine a) Basic principles involved in Ayurveda, Siddha, Unani and			
	Homeopathy			
	b) Preparation and standardization of Ayurvedic formulations			
	viz Aristas and Asawas, Ghutika, Churna, Lehya and Bhasma.			
Unit II	Nutraceuticals	07		
	General aspects, Market, growth, scope and types of products	5		
	available in the market. Health benefits and role of nutraceuticals			
	in ailments like Diabetes, CVS diseases, Cancer, Irritable bowel syndrome and various Gastro intestinal diseases.	1		
	Study of following herbs as health food: Alfaalfa, Chicory,			
	Ginger, Fenugreek, Garlic, Honey, Amla, Ginseng, Ashwagandha,			
	Spirulina			
	Herbal-Drug and Herb-Food Interactions: General			
	introduction to interaction and classification. Study of the	!		
	following drugs and their possible side effects and interactions:			
	Hypericum, kava-kava, Ginkgo Biloba, Ginseng, Garlic, Pepper & Ephedra.			
	Lpneura.			

Unit III	Herbal Cosmetics	10 hours
	Sources and description of raw materials of herbal origin used	10 110413
	via, fixed oils, waxes, gums, colours, perfumes, protective agents,	
	bleaching agents, and antioxidants in products such as skin care,	
	hair care and oral hygiene products.	
	Herbal excipients:	
	Herbal Excipients – Significance of substances of natural origin	
	as excipients – colourants, sweeteners, binders, diluents,	
	viscosity builders, disintegrants, flavours & perfumes.	
	Herbal formulations:	
	Conventional herbal formulations like syrups, mixtures and	
	tablets and Novel dosage forms like phytosomes.	
Unit IV	Evaluation of Drugs WHO & ICH guidelines for the assessment	10 hours
	of herbal drugs stability testing of herbal drugs.	
	Patenting and Regulatory requirements of natural products:	
	a) Definition of the terms: Patent, IPR, Farmers right, Breeder's	
	right, Bioprospecting and	
	Biopiracy	
	b) Patenting aspects of Traditional Knowledge and Natural	
	Products. Case study of Curcuma & Neem. Populatory Jagues - Regulations in India (ASH DTAR ASH DCC)	
	Regulatory Issues - Regulations in India (ASU DTAB, ASU DCC), Regulation of manufacture of ASU drugs - Schedule Z of Drugs &	
	Cosmetics Act for ASU drugs.	
Unit V	General Introduction to Herbal Industry	07
Oiiit v	Herbal drugs industry: Present scope and future prospects. A	hours
	brief account of plant-based industries and institutions involved	nours
	in work on medicinal and aromatic plants in India.	
	Schedule T –Good Manufacturing practice of Indian systems	
	of medicine	
	Components of GMP (Schedule – T) and its objectives	
	Infrastructural requirements, working space, storage area,	
	machinery and equipment, standard operating procedures,	
	health and hygiene, documentation and records.	
	RECOMMENDED BOOKS (LATEST EDITIONS)	

- 1. Textbook of Pharmacognosy by Trease & Evans.
- 2. Textbook of Pharmacognosy by Tyler, Brady & Robber.
- 3. Pharmacognosy by Kokate, Purohit and Gokhale
- 4. Essential of Pharmacognosy by Dr.S.H. Ansari
- 5. Pharmacognosy & Phytochemistry by V.D. Rangari
- 6. Pharmacopoeal standards for Ayurvedic Formulation (Council of Research in Indian Medicine & Homeopathy)
- 7. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation of Botanicals. Business Horizons Publishers, New Delhi, India, 2002.

BP609P	HERBAL DRUG TECHNOLOGY (Practical)	4 Hours/Week
S.No.	Experiment	
1	To perform preliminary phytochemical screening of cru	ide drugs.
2	Determination of the alcohol content of Asava and Arist	a
3	Evaluation of excipients of natural origin	
4	Incorporation of prepared and standardized extract in cosmetic formulations like creams, lotions and shampoos and their evaluation.	
5	Incorporation of prepared and standardized extract in formulations like syrups, mixtures and tablets and their evaluation as per Pharmacopoeial requirements.	
6	Monograph analysis of herbal drugs from recent Pharmacopoeias	
7	Determination of Aldehyde content	
8	Determination of Phenol content	
9	Determination of total alkaloids	

BP604T	BIOPHARMACEUTICS AND PHARMACOKINETICS (THEORY)	45 Hours	
pharmacok	Scope: This subject is designed to impart knowledge and skills of Biopharmaceutics and pharmacokinetics and their applications in pharmaceutical development, design of dose and		
dosage reg	imens and in solving the problems raised therein.		
	COURSE OUTCOMES		
CO 1	Learn about various parameters of drug absorption and distribu		
CO 2	Gain knowledge on drug elimination and metabolism process. Uthe bioavailability and bioequivalence studies.	nderstand about	
CO 3	Understand about the various pharmacokinetic models a pharmacokinetic parameters.	nd analyze the	
CO 4	Locating the various pharmacokinetic parameters in multicomp	artment models.	
CO 5	Assessing the parameters of non-linear pharmacokinetics.		
	COURSE CONTENT		
Unit I	Introduction to Biopharmaceutics Absorption; Mechanisms of drug absorption through GIT	10 hours	
	factors influencing drug absorption though GIT, absorption of	f	
	drug from Non per oral extra-vascular routes.		
	Distribution Tissue permeability of drugs, binding of drugs		
	apparent, volume of drug distribution, plasma and tissue proteir binding of drugs, factors affecting protein-drug binding. Kinetics		
	of protein binding, Clinical significance of protein binding of		
	drugs		
Unit II	Elimination: Drug metabolism and basic understanding	10	
	metabolic pathways renal excretion of drugs, factors affecting	10	
	renal excretion of drugs, renal clearance, Non renal routes of	hours	
	drug excretion of drugs		
	Bioavailability and Bioequivalence: Definition and		
	Objectives of bioavailability, absolute and relative	2	
	bioavailability, measurement of bioavailability, in-vitro drug dissolution models, in-vitro-in-vivo correlations		
	bioequivalence studies, methods to enhance the dissolution		
	rates and bioavailability of poorly soluble drugs.		
Unit III	Pharmacokinetics: Definition and introduction to	10 hours	
	Pharmacokinetics, Compartment models, Non compartment		
	models, physiological models, One compartment open model		
	(a). Intravenous Injection (Bolus) (b). Intravenous infusion and		
	(c)Extravascular administrations. Pharmacokinetics		
	parameters - KE, t1/2, Vd, AUC, Ka, Clt and CL _R - definitions		
	methods of eliminations, understanding of their significance and		
	application		

Unit IV	Multicompartment models: Two compartment open model. IV	10 hours
	bolus. Kinetics of multiple dosing, steady state drug levels,	
	calculation of loading and maintenance doses and their	
	significance in clinical settings.	
Unit V	Nonlinear Pharmacokinetics:	07
	a. Introduction, b. Factors causing non-linearity. c. Michaelis-	hours
	menton method of estimating parameters, Explanation with an	
	example of drugs.	
DECOMMENDED BOOKS (LATEST EDITIONS)		

- 1. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi.
- 2. Biopharmaceutics and Pharmacokinetics; By Robert F Notari
- 3. Applied biopharmaceutics and pharmacokinetics, Leon Shargel and Andrew B.C.YU 4th edition, Prentice-Hall Inernational edition.USA
- 4. Biopharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmankar and Sunil B. Jaiswal, Vallabh Prakashan Pitampura, Delhi
- 5. Pharmacokinetics: ByMilo Glbaldi Donald, R. Mercel Dekker Inc.
- 6. Hand Book of Clinical Pharmacokinetics, ByMilo Gibaldi and Laurie Prescott by ADIS Health Science Press.
- 7. Biopharmaceutics; By Swarbrick
- 8. Clinical Pharmacokinetics, Concepts and Applications: ByMalcolm Rowland and
- 9. Thomas, N. Tozen, Lea and Febrger, Philadelphia, 1995.
- 10. Dissolution, Bioavailability and Bioequivalence, By Abdou H.M, Mack, Publishing Company, Pennsylvania 1989.
- 11. Biopharmaceutics and Clinical Pharmacokinetics-An introduction 4th edition
- 12. Revised and expanded by Rebort F Notari Marcel Dekker Inn, New York and Basel, 1987.
- 13. Remington's Pharmaceutical Sciences, ByMack Publishing Company, Pennsylvnia

		TSEMESTER
BP605T	PHARMACEUTICAL BIOTECHNOLOGY (Theory)	45 Hrs
 Scope: Biotechnology has a long promise to revolutionize the biological sciences and technology. Scientific application of biotechnology in the field of genetic engineering, medicine and fermentation technology makes the subject interesting. Biotechnology is leading to new biological revolutions in diagnosis, prevention and cure of diseases, new and cheaper pharmaceutical drugs. Biotechnology has already produced transgenic crops and animals and the future promises lot more. It is a research-based subject. 		
	COURSE OUTCOMES	
CO 1	Understand the basics of biotechnology, enzyme biotechnol engineering in pharmaceutical applications.	ogy, and genetic
CO 2	Apply genetic engineering techniques, including cloning and r technology, in medical applications.	ecombinant DNA
	Understand immunity types, vaccine production, and the ro	ole of hybridoma

CO 2	Apply genetic engineering techniques, including cloning and rec	combinant DNA
00 -	technology, in medical applications.	
CO 3	Understand immunity types, vaccine production, and the role	of hybridoma
CO 3	technology.	
CO 4	Understand blotting techniques and microbial genetics, including	mutations and
CO 4	transformations.	
COF	Learn fermentation methods, large-scale production process	es, and blood
CO 5	product handling.	
COURSE CONTENT		
** ** *		
Unit I	a) Brief introduction to Biotechnology with reference to	10 hours
	Pharmaceutical Sciences.	

Unit I	a) Brief introduction to Biotechnology with reference to	10 hours
	Pharmaceutical Sciences.	10 Hours
	b) Enzyme Biotechnology- Methods of enzyme immobilization	
	and applications.	
	c) Biosensors- Working and applications of biosensors in	
	Pharmaceutical Industries.	
	d) Brief introduction to Protein Engineering.	
	e) Use of microbes in industry. Production of Enzymes- General	
	consideration -	
	Amylase, Catalase, Peroxidase, Lipase, Protease, Penicillinase.	
	f) Basic principles of genetic engineering.	
Unit II	a) Study of cloning vectors, restriction endonucleases and DNA	10 hours
	ligase.	10 110013
	b) Recombinant DNA technology. Application of genetic	
	engineering in medicine.	
	c) Application of r DNA technology and genetic engineering in	
L		

	the production of:	
	i) Interferon ii) Vaccines- hepatitis- B iii) Hormones-Insulin.	
	d) Brief introduction to PCR	
Unit III	Types of immunity- humoral immunity, cellular immunity	
Omit in	a) Structure of Immunoglobulins	10 hours
	b) Structure and Function of MHC	
	c) Hypersensitivity reactions, Immune stimulation and Immune	
	suppressions.	
	d) General method of the preparation of bacterial vaccines,	
	toxoids, viral vaccine,	
	antitoxins, serum-immune blood derivatives and other products	
	relative to immunity.	
	e) Storage conditions and stability of official vaccines	
	f) Hybridoma technology- Production, Purification and	
	Applications	
11	g) Blood products and Plasma Substitutes.	001
Unit IV	a) Immuno blotting techniques- ELISA, Western blotting, Southern blotting.	08 hours
	b) Genetic organization of Eukaryotes and Prokaryotes	
	c) Microbial genetics including transformation, transduction,	
	conjugation, plasmids and	
	transposons.	
	d) Introduction to Microbial biotransformation and applications.	
	e) Mutation: Types of mutation/mutants.	
Unit V	a) Fermentation methods and general requirements, study of	07 hours
	media, equipment, sterilization methods, aeration process,	
	stirring.	
	b) Large-scale production fermenter design and its various	
	controls.	
	c) Study of the production of - penicillins, citric acid, Vitamin B12, Glutamic acid, Griseofulvin,	
	d) Blood Products: Collection, Processing and Storage of whole	
	human blood, dried human plasma, plasma Substitutes.	
	RECOMMENDED BOOKS (LATEST EDITIONS)	
ALLOCATILITE DO OTTO (MITTED I DETITIONO)		

- 1. Computer Application in Pharmacy William E. Fassett –Lea and Febiger, 600 South Washington Square, USA, (215) 922-1330.
- 2. Computer Application in Pharmaceutical Research and Development –Sean Ekins Wiley-Interscience, A John Willey and Sons, INC., Publication, USA
- 3. Bioinformatics (Concept, Skills and Applications) S.C.Rastogi-CBS Publishers and Distributors, 4596/1- A, 11 Darya Gani, New Delhi 110 002(INDIA)
- 4. Microsoft office Access 2003, Application Development Using VBA, SQL Server, DAP and Infopath Cary N. Prague Wiley Dreamtech India (P) Ltd., 4435/7, Ansari Road, Daryagani, New Delhi 110002

BP606T	PHARMACEUTICAL QUALITY ASSURANCE (Theory)	45 Hrs
Scope: This	uality assurance	
aspects of pharmaceutical industries. It deals with the important aspects like cGMP, QC tests,		
documenta	tion, quality certifications and regulatory affairs.	
	COURSE OUTCOMES	
CO 1	Understand the principles of quality management systems in the	pharmaceutical
COI	industry.	
CO 2	Analyze the regulatory requirements and guidelines for pharma	aceutical quality
	assurance.	
CO 3	Evaluate the processes and procedures for Good Manufacturing I	Practices (GMP).
CO 4	Explain the principles of validation and qualification in	pharmac eutical
60 4	manufacturing.	
CO 5	Apply quality assurance principles to ensure product quality and	documentations
000	for patient safety.	
	COURSE CONTENT	T
Unit I	Quality Assurance and Quality Management concepts:	TO HOULS
	Definition and concept of Quality control, Quality assurance and	
	GMP.	
	Total Quality Management (TQM): Definition, elements, philosophies.	
	ICH Guidelines: purpose, participants, process of	
	harmonization, Brief overview of QSEM, with special emphasis	
	on Q-series guidelines, ICH stability testing guidelines	
	Quality by design (QbD): Definition, overview, elements of	
	QbD program, tools.	
	ISO 9000 & ISO14000: Overview, Benefits, Elements, steps for	
	registration	
	NABL accreditation: Principles and procedures	
Unit II	Organization and personnel: Personnel responsibilities,	10 hours
	training, hygiene and personal records.	
	Premises: Design, construction and plant layout, maintenance,	
	sanitation, environmental control, utilities and maintenance of	
	sterile areas, control of contamination.	
	Equipment and raw materials: Equipment selection,	
	purchase specifications, maintenance, purchase specifications and maintenance of stores for raw materials.	
Unit III	Quality Control: Quality control test for containers, rubber	
Omit III	closures and secondary packing materials.	10 hours
	Good Laboratory Practices: General Provisions, Organization	
	and Personnel, Facilities, Equipment, Testing Facilities Operation, Test and Control Articles, Protocol for Conduct of a	

	Nonclinical Laboratory Study, Records and Reports,		
	Disqualification of Testing Facilities		
Unit IV	Complaints: Complaints and evaluation of complaints, Handling	08 hours	
	of return good, recalling and waste disposal.	oonours	
	Document maintenance in pharmaceutical industry: Batch		
	Formula Record, Master Formula Record, SOP, Quality audit,		
	Quality Review and Quality documentation, Reports and		
	documents, distribution records.		
Unit V	Calibration and Validation: Introduction, definition and	07 hours	
	general principles of calibration, qualification and validation,		
	importance and scope of validation, types of validation, and		
	validation master plan. Calibration of pH meter, Qualification of		
	UV-visible spectrophotometer, General principles of Analytical		
	method Validation.		
	Warehousing: Good warehousing practice, materials		
	management		
DECOMMENDED DOOVE (LATECT EDITIONE)			

- 1. Quality Assurance Guide by organization of Pharmaceutical Products of India.
- 2. Good Laboratory Practice Regulations, 2nd Edition, Sandy Weinberg Vol. 69.
- 3. Quality Assurance of Pharmaceuticals- A compendium of Guidelines and Related materials Vol I WHO Publications.
- 4. A guide to Total Quality Management- Kushik Maitra and Sedhan K Ghosh
- 5. How to Practice GMP's P Sedan Sharma.
- 6. ISO 9000 and Total QualityManagement Sadhank G Ghosh
- 7. The International Pharmacopoeia Vol I, II, III, IV- General Methods of Analysis and Quality specification for Pharmaceutical Substances, Excipients and Dosage forms
- 8. Good laboratory Practices Marcel Deckker Series
- 9. ICH guidelines, ISO 9000 and 14000 guidelines

	B. PHARM (SEMESTER – VII)		
BP701T	INSTRUMENTAL METHODS OF ANALYSIS (Theory)	45 Hours	
quantitativ on the prir This also	Scope: This subject deals with the application of instrumental methods in qualitative an quantitative analysis of drugs. This subject is designed to impart a fundamental knowledg on the principles and instrumentation of spectroscopic and chromatographic technique. This also emphasizes on theoretical and practical knowledge on modern analytical instruments that are used for drug testing.		
	COURSE OUTCOMES		
CO 1	Understand the principles and instrumentation of various techniques.	analytical	
CO 2	Analyze the applications of spectroscopic methods in pharmaceutic	cal analysis.	
CO 3	Evaluate the principles and techniques of chromatographic separation.		
CO 4	Explain the principles of sophisticated chromatographic analysis and its applications.		
CO 5	Apply instrumental methods in the quantitative and qualitative analysis of drugs.		
	COURSE CONTENT		
Unit I	UV Visible spectroscopy Electronic transitions, chromophores, auxochromes, spectral shifts, solvent effect on absorption spectra, Beer and Lambert's law, Derivation and deviations. Instrumentation - Sources of radiation, wavelength selectors, sample cells, detectors- Photo tube, Photomultiplier tube, Photo voltaic cell, Silicon Photodiode. Applications - Spectrophotometric titrations, Single component and multi component analysis. Fluorimetry Theory, Concepts of singlet, doublet and triplet electronic states, internal and external conversions, factors affecting fluorescence, quenching, instrumentation and applications	hours	
Unit II	IR spectroscopy Introduction fundamental modes of vibrations in polyatomic	10	

Introduction, fundamental modes of vibrations in polyatomic

Instrumentation - Sources of radiation, wavelength selectors, detectors - Golay cell, Bolometer, Thermocouple, Thermistor,

Flame Photometry-Principle, interferences, instrumentation and

molecules, sample handling, factors affecting vibrations.

Pyroelectric detector and applications.

hours

Unit III	Atomic absorption spectroscopy- Principle, interferences, instrumentation and applications. Nepheloturbidometry- Principle, instrumentation, and applications. Introduction to chromatography Adsorption and partition column chromatography- Methodology, advantages, disadvantages and applications. Thin layer chromatography- Introduction, Principle, Methodology, Rf values, advantages, disadvantages and applications. Paper chromatography-Introduction, methodology, development techniques, advantages, disadvantages and applications. Electrophoresis- Introduction, factors affecting electrophoretic	10 hours
Unit IV	mobility, Techniques of paper, gel, capillary electrophoresis, applications Gas chromatography- Introduction, theory, instrumentation, derivatization, temperature programming, advantages, disadvantages and applications. High-performance liquid chromatography (HPLC)	08 hours
Unit V	Introduction, theory, instrumentation, advantages and applications. Ion exchange chromatography- Introduction, classification, ion exchange resins, properties, mechanism of ion exchange process, factors affecting ion exchange, methodology and applications. Gel chromatography- Introduction, theory, instrumentation and applications. Affinity chromatography- Introduction, theory, instrumentation	07 hours

- 1. Instrumental Methods of Chemical Analysis by B.K Sharma
- 2. Organic spectroscopy by Y.R Sharma
- 3. Textbook of Pharmaceutical Analysis by Kenneth A. Connors
- 4. Vogel's Textbook of Quantitative Chemical Analysis by A.I. Vogel
- 5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake
- 6. Organic Chemistry by I. L. Finar
- 7. Organic spectroscopy by William Kemp
- 8. Quantitative Analysis of Drugs by D. C. Garrett
- 9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi
- 10. Spectrophotometric identification of Organic Compounds by Silverstein

BP705P	INSTRUMENTAL METHODS OF ANALYSIS (Practical)	4 Hours/Week
S.No.	Topic	
1.	Determination of absorption maxima and effect of solv	vents absorption
	maxima of organic compounds	
2.	Estimation of dextrose by colorimetry	
3.	Estimation of sulfanilamide by colorimetry	
4.	Simultaneous estimation of ibuprofen and paracetamol by U	IV spectroscopy
5.	Assay of paracetamol by UV- Spectrophotometry	
6.	Estimation of quinine sulfate by fluorimetry	
7.	Study of quenching of fluorescence	
8.	Determination of sodium by flame photometry	
9.	Determination of potassium by flame photometry	
10.	Determination of chlorides and sulphates by Nepheloturbidometry	
11.	Separation of amino acids by paper chromatography	
12.	Separation of sugars by thin layer chromatography	
13.	Separation of plant pigments by column chromatography	
14.	Demonstration experiment on HPLC	
15.	Demonstration experiment on Gas Chromatography	

BP702T	INDUSTRIAL PHARMACY-II (Theory)	45 Hours	
Scope: This course is designed to impart fundamental knowledge on pharmac		ceutical product	
development and translation from laboratory to market.			
COURSE OUTCOMES			
CO 1	Analyze the significance of personnel, space, and raw material requirements in pilot plant scale-up processes.		
CO 2	Demonstrate the ability to apply WHO guidelines for technology transfer		
CO 3	Evaluate the historical evolution and current role of regulatory affairs		
CO 4	Differentiate between various quality management concents such as OhD and Six		
CO 5	Explaining the responsibilities of CDSCO and State Licensing interpreting regulatory requirements for new drugs in India.	Authorities and	
	COURSE CONTENT		
Unit I	Pilot plant scale up techniques: General considerations -	TO HOURS	
	including significance of personnel requirements, space		
	requirements, raw materials, Pilot plant scale-up considerations		
	for solids, liquid orals, semi-solids and relevant documentation,		
IInit II	SUPAC guidelines, Introduction to platform technology.		
Unit II	Technology development and transfer: WHO guidelines for Technology Transfer(TT):	10 hours	
	Terminology, Technology transfer protocol, Quality risk		
	management, Transfer from R & D to production (Process,		
	packaging and cleaning), Granularity of TT Process (API,		
	excipients, finished products, packaging materials)		
	Documentation, Premises and equipment, qualification and		
	validation, quality control, analytical method transfer,		
	Approved regulatory bodies and agencies, Commercialization -		
	practical aspects and problems (case studies), TT agencies in		
	India - APCTD, NRDC, TIFAC, BCIL, TBSE / SIDBI; TT related		
	documentation - confidentiality agreement, licensing, MoUs,		
	legal issues.		
Unit III	Regulatory affairs : Introduction, Historical overview of	10 110013	
	Regulatory Affairs, Regulatory authorities, Role of Regulatory		
	affairs department, Responsibility of Regulatory Affairs Professionals.		
	Regulatory requirements for drug approval: Drug		
	Development Teams, Non-Clinical Drug Development,		
	r	ı	

	Pharmacology, Drug Metabolism and Toxicology, General	
	considerations of Investigational New Drug (IND) Application,	
	Investigator's Brochure (IB) and New Drug Application (NDA),	
	Clinical research / BE studies, Clinical Research Protocols,	
	Biostatistics in Pharmaceutical Product Development, Data	
	Presentation for FDA Submissions, Management of Clinical	
	Studies.	
Unit IV	Quality management systems: Quality management &	08 hours
	Certifications: Concept of Quality, Total Quality Management,	
	Quality by Design (QbD), Six Sigma concept, Out of Specifications	
	(OOS), Change control, Introduction to ISO 9000 series of quality	
	systems standards, ISO 14000, NABL, GLP.	
Unit V	Indian Regulatory Requirements: Central Drug Standard	07 hours
	Control Organization (CDSCO) and State Licensing Authority:	
	Organization, Responsibilities, Certificate of Pharmaceutical	
	Product (COPP), Regulatory requirements and approval	
	procedures for New Drugs.	

- Regulatory Affairs from Wikipedia, the free encyclopedia modified on 7th April 1. available at http://en.wikipedia.org/wiki/Regulatory_Affairs.
- International Regulatory 2. **Affairs** Updates, 2005. available at http://www.iraup.com/about.php
- Douglas J Pisano and David S. Mantus. Textbook of FDA Regulatory Affairs A Guide 3. for Prescription Drugs, Medical Devices, and Biologics' Second Edition.
- Affairs brought by Learning Plus, inc. 4. Regulatory available at http.//www.cgmp.com/ra.htm.

BP703T	PHARMACY PRACTICE (Theory)	45 Hours
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Scope: In the changing scenario of pharmacy practice in India, for successful practice of Hospital Pharmacy, the students are required to learn various skills like drug distribution, drug information, and therapeutic drug monitoring for improved patient care. In community pharmacy, students will be learning various skills such as dispensing of drugs, responding to minor ailments by providing suitable safe medication, and patient counselling for improved patient care in the community set up.

improved patient care in the community set up.		
COURSE OUTCOMES		
CO 1	Understand hospital structures, the role of hospital pharmacies adverse drug reactions effectively.	, and manage
CO 2	Implement drug distribution systems, maintain hospital formularies, and enhance medication adherence and community pharmacy management.	
CO 3	Manage pharmacy committees, provide drug information, and co effectively.	unsel patients
CO 4	Prepare hospital budgets, perform clinical pharmacy functions, rational OTC medication use.	and promote
CO 5	Manage drug inventory, understand investigational drugs, and int laboratory tests.	erpret clinical
	COURSE CONTENT	
Unit I	a) Hospital and it's organization Definition, Classification of hospital- Primary, Secondary and Tertiary hospitals, Classification based on clinical and non-clinical basis, Organization Structure of a Hospital, and Medical staffs involved in the hospital and their functions. b) Hospital pharmacy and its organization Definition, functions of hospital pharmacy, Organization structure, Location, Layout and staff requirements, and Responsibilities and functions of hospital pharmacists. c) Adverse drug reaction Classifications - Excessive pharmacological effects, secondary pharmacological effects, idiosyncrasy, allergic drug reactions, genetically determined toxicity, toxicity following the sudden withdrawal of drugs, Drug interaction-beneficial interactions, adverse interactions, and pharmacokinetic drug interactions, Methods for detecting drug interactions, spontaneous case reports and record linkage studies, and Adverse drug reaction reporting and	10 hours
Unit II	management. a) Drug distribution system in a hospital	10 hours

Dispensing of drugs to inpatients, types of drug distribution systems, charging policy and labelling, dispensing of drugs to ambulatory patients, and Dispensing of controlled drugs.

b) Hospital formulary

Definition, contents of hospital formulary, Differentiation of hospital formulary and Drug list, preparation and revision, and addition and deletion of drug from hospital formulary.

c) Therapeutic drug monitoring

Need for Therapeutic Drug Monitoring, Factors to be considered during the Therapeutic Drug Monitoring, and Indian scenario for Therapeutic Drug Monitoring.

d) Medication adherence

Causes of medication non-adherence, pharmacist role in medication adherence, and monitoring of patient medication adherence.

e) Patient medication history interview

Need for the patient medication history interview, medication interview forms.

f) Community pharmacy management

Financial, materials, staff, and infrastructure requirements.

Unit III

a) Pharmacy and therapeutic committee

Organization, functions, and Policies of the pharmacy and therapeutics committee in including drugs in the formulary, inpatient and outpatient prescriptions, automatic stop order, and emergency drug list preparation.

b) Drug information services

Drug and Poison information centre, Sources of drug information, Computerized services, and storage and retrieval of information.

c) Patient counselling:

Definition of patient counselling; steps involved in patient counselling, and Special cases that require the pharmacist

d) Education and training program in the hospital

Role of pharmacist in the education and training program, Internal and external training program, Services to the nursing homes/clinics, Code of ethics for community pharmacy, and Role of pharmacist in the interdepartmental communication and community health education.

e) Prescribed medication order and communication skills

Prescribed medication order- interpretation and legal requirements, and Communication skills- communication with prescribers and patients.

Unit IV

a) Budget preparation and implementation

10 hours

10 hours

	b) Clinical Pharmacy	
	Introduction to Clinical Pharmacy, Concept of clinical pharmacy,	
	functions and responsibilities of clinical pharmacist, Drug	
	therapy monitoring - medication chart review, clinical review,	
	pharmacist intervention, Ward round participation, Medication	
	history and pharmaceutical care.	
	Dosing pattern and drug therapy based on Pharmacokinetic &	
	disease pattern.	
	c) Over the counter (OTC) sales	
	Introduction and sale of over the counter, and Rational use of	
	common over the counter medications.	
Unit V	a) Drug store management and inventory control	07
Unit V	a) Drug store management and inventory control Organization of drug store, types of materials stocked and	07 hours
Unit V		_
Unit V	Organization of drug store, types of materials stocked and	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles,	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking,	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking, Economic order quantity, Reorder quantity level, and Methods	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking, Economic order quantity, Reorder quantity level, and Methods used for the analysis of the drug expenditure	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking, Economic order quantity, Reorder quantity level, and Methods used for the analysis of the drug expenditure b) Investigational use of drugs	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking, Economic order quantity, Reorder quantity level, and Methods used for the analysis of the drug expenditure b) Investigational use of drugs Description, principles involved, classification, control, identification, role of a hospital pharmacist, advisory committee.	_
Unit V	Organization of drug store, types of materials stocked and storage conditions, Purchase and inventory control: principles, purchase procedure, purchase order, procurement and stocking, Economic order quantity, Reorder quantity level, and Methods used for the analysis of the drug expenditure b) Investigational use of drugs Description, principles involved, classification, control,	_

- 1. Merchant S.H. and Dr. J.S.Quadry. *A textbook of hospital pharmacy*, 4th ed. Ahmadabad: B.S. Shah Prakakshan; 2001.
- 2. Parthasarathi G, Karin Nyfort-Hansen, Milap C Nahata. *A textbook of Clinical Pharmacy Practice- essential concepts and skills,* 1st ed. Chennai: Orient Longman Private Limited; 2004.
- 3. William E. Hassan. *Hospital pharmacy*, 5th ed. Philadelphia: Lea & Febiger;1986.
- 4. Tipnis Bajaj. *Hospital Pharmacy*, 1st ed. Maharashtra: Career Publications; 2008.
- 5. Scott LT. *Basic skills in interpreting laboratory data*, 4thed. American Society of Health System Pharmacists Inc; 2009.
- 6. Parmar N.S. *Health Education and Community Pharmacy,* 18th ed. India: CBS Publishers & Distributers; 2008.

Journals:

- 1. Therapeutic drug monitoring. ISSN: 0163-4356
- 2. Journal of pharmacy practice. ISSN: 0974-8326
- 3. American journal of health system pharmacy. ISSN: 1535-2900 (online)
- 4. Pharmacy times (Monthly magazine)

	NOVEL DRIVE DELIVEDIV GUERRIA (TIVE DRIV	45 Hours	
BP704T	NOVEL DRUG DELIVERY SYSTEMS (THEORY)		
Scope: This subject is designed to impart basic knowledge on the area of novel		el drug delivery	
systems.			
	COURSE OUTCOMES	. 11 1 1	
CO 1	Classify and justify the selection of drug candidates for corformulations using diffusion, dissolution, and ion exchange princ		
	Evaluate the advantages and disadvantages of microencapsulation and mucos		
CO 2	drug delivery systems.		
CO 3	Compare and contrast different transdermal and gastro retentive drug delivery systems, including formulation approaches and applications.		
CO 4	Assess the advantages and disadvantages of targeted drug delivery systems such		
CO 4	as liposomes, niosomes, nanoparticles, and monoclonal antibodic		
	Analyze intraocular barriers and methods to overcome them	_	
CO 5	delivery systems, and evaluate the advantages and disadvantages	s of intrauterine	
	drug delivery systems. COURSE CONTENT		
	COURSE CONTENT		
Unit I	Controlled drug delivery systems: Introduction, terminology	10 hours	
	or definitions and rationale, advantages, disadvantages, selection	10 Hours	
	of drug candidates. Approaches to design controlled release		
	formulations based on diffusion, dissolution and ion exchange		
	principles. Physicochemical and biological properties of drugs		
	relevant to controlled release formulations.		
	Polymers: Introduction, classification, properties, advantages		
	and application of polymers in formulation of controlled release		
** ** **	drug delivery systems.		
Unit II	Microencapsulation: Definition, advantages and disadvantages,	10	
	microspheres /microcapsules, microparticles, methods of	hours	
	microencapsulation, applications. Mucosal Drug Delivery system: Introduction, Principles of bio	nours	
	adhesion/mucoadhesion, concepts, advantages and		
	disadvantages, transmucosal permeability and formulation		
	considerations of buccal delivery systems.		
	Implantable Drug Delivery Systems: Introduction, advantages		
	and disadvantages, concept of implants and osmotic pump		
Unit III	Transdermal Drug Delivery Systems: Introduction,	10 hours	
	Permeation through skin, factors affecting permeation,		
	permeation enhancers, basic components of TDDS, formulation		
	approaches.		
	Gastro retentive drug delivery systems: Introduction,		
	advantages, disadvantages, approaches for GRDDS – Floating,		
	high-density systems, inflatable and gastro adhesive systems		

	and their applications	
	Nasopulmonary drug delivery system: Introduction to Nasal	
	and Pulmonary routes of drug delivery, Formulation of Inhalers	
	(dry powder and metered dose), nasal sprays, nebulizers.	
Unit IV	Targeted drug Delivery: Concepts and approaches advantages	08 hours
	and disadvantages, introduction to liposomes, niosomes,	
	nanoparticles, monoclonal antibodies and their applications	
Unit V	Ocular Drug Delivery Systems: Introduction, intra ocular	07 hours
	barriers and methods to overcome –Preliminary study, ocular	
	formulations and ocuserts.	
	Intrauterine Drug Delivery Systems: Introduction, advantages	
	and disadvantages, development of intrauterine devices (IUDs)	
	and applications	

- 1. Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded, Marcel Dekker, Inc., New York, 1992.
- 2. Robinson, J. R., Lee V. H. L, Controlled Drug Delivery Systems, Marcel Dekker, Inc., New York, 1992.
- 3. Encyclopedia of Controlled Delivery. Edith Mathiowitz, Published by Wiley Interscience Publication, John Wiley and Sons, Inc, New York. Chichester/Weinheim
- 4. N.K. Jain, Controlled and Novel Drug Delivery, CBS Publishers & Distributors, New Delhi, First edition 1997 (reprint in 2001).
- 5. S.P. Vyas and R.K. Khar, Controlled Drug Delivery -concepts and advances, Vallabh Prakashan, New Delhi, First edition 2002.

Journals

- 6. Indian Journal of Pharmaceutical Sciences (IPA)
- 7. Indian Drugs (IDMA)
- 8. Journal of Controlled Release (Elsevier Sciences)
- 9. Drug Development and Industrial Pharmacy (Marcel & Decker)
- 10. International Journal of Pharmaceutics (Elsevier Sciences)

B. PHARM (SEMESTER - VIII)	
BIOSTATISITCS AND RESEARCH METHODOLOGY (Theory)	45 Hours

Scope: To understand the applications of Biostatics in Pharmacy. This subject deals with descriptive statistics, Graphics, Correlation, Regression, logistic regression Probability theory, Sampling technique, Parametric tests, Non-Parametric tests, ANOVA, Introduction to Design of Experiments, Phases of Clinical trials and Observational and Experimental studies, SPSS, R and MINITAB statistical software's, analyzing the statistical data using Excel.

BP801T

COURSE OUTCOMES		
CO 1	Explain and apply measures of central tendency and measures of dusing pharmaceutical examples.	ispersion
CO 2	Analyze pharmaceutical data using regression techniques and probability distributions.	
CO 3	Evaluate the appropriateness of non-parametric tests in pharmaceutical research, and design experiments using graphical representations.	
CO 4	Assess blocking and confounding in factorial designs, conduct hypothesis testing in regression models, and apply statistical analysis using tools.	
CO 5	CO 5 Appraise factorial designs and response surface methodology for pharmaceutical research	
COURSE CONTENT		
Unit I	Introduction: Statistics, Biostatistics, Frequency distribution	10
	Measures of central tendency: Mean, Median, Mode-	hours
	Pharmaceutical examples	
	Measures of dispersion: Dispersion, Range, standard deviation,	
	pharmaceutical problems Correlation: Definition, Karl Pearson's coefficient of correlation,	
	Multiple correlation -Pharmaceuticals examples	
Unit II	Regression: Curve fitting by the method of least squares, fitting	10
	the lines $y = a + bx$ and $x = a + by$, Multiple regression, standard error	10
	of regression – Pharmaceutical Examples	hours
	Probability: Definition of probability, Binomial distribution,	
	Normal distribution, Poisson's distribution, properties - problems	
	Sample, Population, large sample, small sample, Null hypothesis,	
	alternative hypothesis, sampling, essence of sampling, types of	
	sampling, Error-I type, Error-II type, Standard error of mean (SEM) - Pharmaceutical examples	

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- 1. Pharmaceutical statistics- Practical and clinical applications, Sanford Bolton, publisher Marcel Dekker Inc. New York.
- 2. Fundamental of Statistics Himalaya Publishing House- S.C. Gupta
- 3. Design and Analysis of Experiments -PHI Learning Private Limited, R. Panneerselvam,
- 4. Design and Analysis of Experiments Wiley Students Edition, Douglas and C. Montgomery

BP802T	SOCIAL AND PREVENTIVE PHARMACY (Theory)	45 Hours	
Scope: The	purpose of this course is to introduce to students a number of h	ealth issues and	
their challe	enges. This course also introduced a number of national health pr	ogrammes. The	
roles of the	roles of the pharmacist in these contexts are also discussed.		
	COURSE OUTCOMES		
60.4	Understand health and disease concepts, including public heal	th, nutrition, and	
CO 1	socio-cultural health factors.		
CO 2	Apply prevention and control methods for infectious and chronic diseases.		
CO 3	Understand key national health programs and their impact.		
CO 4	Evaluate health interventions for specific populations and WHO	's role.	
CO 5	Recognize the role of community health services and education		
	, ,		
	COURSE CONTENT		
Unit I	Concept of health and disease: Definition, concepts and	1 401	
	evaluation of public health. Understanding the concept o	10 Hours	
	prevention and control of disease, social causes of diseases and		
	social problems of the sick.		
	Social and health education : Food in relation to nutrition and	1	
	health, Balanced diet, Nutritional deficiencies, Vitamin		
	deficiencies, Malnutrition and its prevention.		
	Sociology and health : Socio-cultural factors related to health	,	
	and disease, Impact of urbanization on health and disease		
	Poverty and health.	'	
	Hygiene and health: personal hygiene and health care		
	avoidable habits	,	
Unit II		1	
Uniti	Preventive medicine: General principles of prevention and control of diseases such as cholera, SARS, Ebola virus, influenza	10 Hours	
		·	
	acute respiratory infections, malaria, chicken guinea, dengue		
	lymphatic filariasis, pneumonia, hypertension, diabetes mellitus	,	
Unit III	cancer, drug addiction and drug substance abuse.	1	
Unitin	National health programs, its objectives, functioning and	10 hours	
	outcome of the following:		
	HIV AND AIDS control program, TB, Integrated disease		
	surveillance program (IDSP), National leprosy control program		
	National mental health program, National program for		
	prevention and control of deafness, Universal immunization		
	program, National program for control of blindness, Pulse polic)	
	program.		
Unit IV	National health intervention program for mother and child		

National family welfare program, National tobacco control

program, National Malaria Prevention Program, National	
program for the health care for the elderly, social health program;	
role of WHO in Indian national program.	
Community services in rural, urban and school health: Functions	07 hours
of PHC, Improvement in rural sanitation, national urban health	
mission, Health promotion and education in school.	
DECOMMENDED DOOVE (LATECT EDITIONE)	
hort Textbook of Preventive and Social Medicine, Prabhakara Gl	N, 2nd Edition,
010, ISBN: 9789380704104, JAYPEE Publications	
'extbook of Preventive and Social Medicine (Mahajan and Gupta),	Edited by Roy
abindra Nath, Saha Indranil, 4th Edition, 2013, ISBN: 978935090	01878, JAYPEE
ublications	
Review of Preventive and Social Medicine (Including Biostatistics)	, Jain Vivek, 6 th
dition, 2014, ISBN: 9789351522331, JAYPEE Publications	
ssentials of Community Medicine—A Practical Approach, Hire	math Lalita D.
	-,,,
	on 2011 ISBN-
4: 9788190128285, BANARSIDAS BHANOT PUBLISHERS.	JII, 2011, 10DI
	program for the health care for the elderly, social health program; role of WHO in Indian national program. Community services in rural, urban and school health: Functions of PHC, Improvement in rural sanitation, national urban health mission, Health promotion and education in school. RECOMMENDED BOOKS (LATEST EDITIONS) hort Textbook of Preventive and Social Medicine, Prabhakara Gl 010, ISBN: 9789380704104, JAYPEE Publications extbook of Preventive and Social Medicine (Mahajan and Gupta), tabindra Nath, Saha Indranil, 4th Edition, 2013, ISBN: 978935096 (albications) eview of Preventive and Social Medicine (Including Biostatistics) (dition, 2014, ISBN: 9789351522331, JAYPEE Publications) essentials of Community Medicine—A Practical Approach, Hire diremath Dhananjaya A, 2nd Edition, 2012, ISBN: 978935025 (albications) eark Textbook of Preventive and Social Medicine, K Park, 21st Edition

Community Pharmacy Practice, Ramesh Adepu, BSP publishers, Hyderabad

Research in Social and Administrative Pharmacy, Elsevier, Ireland

6.

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Recommended Journals:

BP803ET	PHARMA MARKETING MANAGEMENT (Theory)	45 Hours

Scope: The pharmaceutical industry not only needs highly qualified researchers, chemists and technical people, but also requires skilled managers who can take the industry forward by managing and taking the complex decisions which are imperative for the growth of the industry. The Knowledge and Know-how of marketing management groom the people for taking a challenging role in Sales and Product management.

taking a challenging role in Sales and Product management.		
COURSE OUTCOMES		
CO 1	Differentiate between marketing and selling, analyze the environment, and evaluate consumer buying behavior in the planter.	_
CO 2	Apply product classification, product life cycle, and product strategies in the pharmaceutical industry.	t positioning
CO 3	Justify the selection of promotional methods, determine promotional assess the effectiveness of various promotional technical sector.	_
CO 4	Formulate pharmaceutical marketing channel strategies, man distribution effectively, and evaluate the roles and responsibilities	
CO 5	Assess pricing methods, strategies, and issues in price management pharmaceutical industry, and analyze emerging concepts like vertical and global marketing.	
	COURSE CONTENT	
Unit I	Marketing:	10 hours
	Definition, general concepts and scope of marketing; Distinction	20 110 410
	between marketing & selling; Marketing environment; Industry	
	and competitive analysis; Analyzing consumer buying behavior; industrial buying behaviour.	
	Pharmaceutical market:	
	Quantitative and qualitative aspects; size and composition of the	
	market; demographic descriptions and socio-psychological	
	characteristics of the consumer; market segmentation&	
	targeting. Consumer profile; Motivation and prescribing habits of the physician; patients' choice of physician and retail	
	pharmacist. Analyzing the Market; Role of market research.	
Unit II	Product decision:	10 h s
	Classification, product line and product mix decisions, product	10 hours
	life cycle, product portfolio analysis; product positioning; New	
	product decisions; Product branding, packaging and labelling	
	decisions, Product management in the pharmaceutical industry.	
Unit III	Promotion:	10 hours

	Methods, determinants of promotional mix, promotional budget;	
	An overview of personal selling, advertising, direct mail,	
	journals, sampling, retailing, medical exhibition, public relations,	
	online promotional techniques for OTC Products.	
Unit IV	Pharmaceutical marketing channels:	08 hours
	Designing channels, channel members, selecting the appropriate	
	channel, conflict in channels, physical distribution management:	
	Strategic importance, tasks in physical distribution management.	
	Professional sales representative (PSR):	
	Duties of PSR, purpose of detailing, selection and training,	
	supervising, norms for customer calls, motivating, evaluating,	
	compensation and future prospects of the PSR.	
Unit V	Pricing:	07
	Meaning, importance, objectives, determinants of price; pricing	hours
	methods and strategies, issues in price management in	
	pharmaceutical industry. An overview of DPCO (Drug Price	
	Control Order) and NPPA (National Pharmaceutical Pricing	
	Authority).	
	Emerging concepts in marketing:	
	Vertical & Horizontal Marketing; Rural Marketing; Consumerism;	
	Industrial Marketing; Global Marketing.	

- 1. Philip Kotler and Kevin Lane Keller: Marketing Management, Prentice Hall of India, New Delhi
- 2. Walker, Boyd and Larreche: Marketing Strategy- Planning and Implementation, Tata MC GrawHill, New Delhi.
- 3. Dhruv Grewal and Michael Levy: Marketing, Tata MC Graw Hill
- 4. Arun Kumar and N Menakshi: Marketing Management, Vikas Publishing, India
- 5. Rajan Saxena: Marketing Management; Tata MC Graw-Hill (India Edition)
- 6. Ramaswamy, U.S & Nanakamari, S: Marketing Management: Global Perspective, Indian Context, Macmilan India, New Delhi.
- 7. Shanker, Ravi: Service Marketing, Excell Books, New Delhi
- 8. Subba Rao Changanti, Pharmaceutical Marketing in India (GIFT Excel series) Excel Publications.

45 Hours PHARMACEUTICAL REGULATORY SCIENCE (THEORY) **BP804ET**

Scope: This course is designed to impart the fundamental knowledge on the regulatory requirements for approval of new drugs, and drug products in regulated markets of India & other countries like the US, EU, Japan, Australia, UK etc. It prepares the students to learn in detail on the regulatory requirements, documentation requirements, and registration procedures for marketing the drug products

procedures for marketing the drug products.			
	COURSE OUTCOMES		
CO 1	Outline the stages of drug discovery and development, include studies and clinical trials.	ing pre-clinical	
CO 2	Analyze the regulatory approval processes for Investigational No New Drug Application (NDA), and Abbreviated New Drug Applica		
CO 3	Explain the procedures for exporting pharmaceutical products, products, products and common like Drug Master Files (DMF) and Common Technic (CTD), and understand international regulatory requirements such ACTD.	ical Documents	
CO 4	Formulate clinical trial protocols, understand the roles of Instit Boards (IRBs) and Independent Ethics Committees, pharmacovigilance and safety monitoring in clinical trials.	and manage	
CO 5	Differentiate between various regulatory terminologies, guidel such as those found in the Orange Book, Federal Register, and of documents.		
COURSE CONTENT			
Unit I	New Drug Discovery and development	10 hours	
	Stages of drug discovery, Drug development process, pre-clinical	10110415	
	studies, non-clinical activities, clinical studies, Innovator and		
	generics, Concept of generics, Generic drug product		
77 '- 77	development.		
Unit II	Regulatory Approval Process Approval processes and timelines involved in Investigational	10	
	New Drug (IND), New Drug Application (NDA), Abbreviated New	hours	
	Drug Application (ANDA), Changes to an approved NDA / ANDA.		
	Drug Application (ANDA). Changes to an approved NDA / ANDA. Regulatory authorities and agencies		
	Drug Application (ANDA). Changes to an approved NDA / ANDA. Regulatory authorities and agencies Overview of regulatory authorities of India, United States,		
	Regulatory authorities and agencies		
	Regulatory authorities and agencies Overview of regulatory authorities of India, United States, European Union, Australia, Japan, Canada (Organization structure and types of applications)		
Unit III	Regulatory authorities and agencies Overview of regulatory authorities of India, United States, European Union, Australia, Japan, Canada (Organization structure and types of applications) Registration of Indian drug product in overseas market	10 hours	
Unit III	Regulatory authorities and agencies Overview of regulatory authorities of India, United States, European Union, Australia, Japan, Canada (Organization structure and types of applications) Registration of Indian drug product in overseas market Procedure for export of pharmaceutical products, Technical	10 hours	
Unit III	Regulatory authorities and agencies Overview of regulatory authorities of India, United States, European Union, Australia, Japan, Canada (Organization structure and types of applications) Registration of Indian drug product in overseas market Procedure for export of pharmaceutical products, Technical documentation, Drug Master Files (DMF), Common Technical	10 hours	
Unit III	Regulatory authorities and agencies Overview of regulatory authorities of India, United States, European Union, Australia, Japan, Canada (Organization structure and types of applications) Registration of Indian drug product in overseas market Procedure for export of pharmaceutical products, Technical	10 hours	

Unit IV	Clinical trials	08 hours
	Developing clinical trial protocols, Institutional Review Board /	
	Independent Ethics committee - formation and working	
	procedures, Informed consent process and procedures, GCP	
	obligations of Investigators, sponsors & Monitors, Managing and	
	Monitoring clinical trials, Pharmacovigilance - safety monitoring	
	in clinical trials	
Unit V	Regulatory Concepts	07 hours
	Basic terminology, guidance, guidelines, regulations, Laws and	
	Acts, Orange book, Federal Register, Code of Federal Regulatory,	
	Purple book	
DECOMMENDED BOOKS (LATEST EDITIONS)		

- 1. Drug Regulatory Affairs by Sachin Itkar, Dr. N.S. Vyawahare, Nirali Prakashan.
- 2. The Pharmaceutical Regulatory Process, Second Edition Edited by Ira R. Berry and Robert P. Martin, Drugs and the Pharmaceutical Sciences, Vol.185. Informa Health care Publishers.
- 3. New Drug Approval Process: Accelerating Global Registrations by Richard A Guarino, MD, 5th edition, Drugs and the Pharmaceutical Sciences, Vol.190.
- 4. Guidebook for drug regulatory submissions / Sandy Weinberg. By John Wiley & Sons. Inc.
- 5. FDA Regulatory Affairs: a guide for prescription drugs, medical devices, and biologics /edited by Douglas J. Pisano, David Mantus.
- 6. Generic Drug Product Development, Solid Oral Dosage forms, Leon Shargel and Isader Kaufer, Marcel Dekker series, Vol.143
- 7. Clinical Trials and Human Research: A Practical Guide to Regulatory Compliance by Fay A. Rozovsky and Rodney K. Adams
- 8. Principles and Practices of Clinical Research, Second Edition Edited by John I. Gallin and Frederick P. Ognibene
- 9. Drugs: From Discovery to Approval, Second Edition by Rick Ng

	VII	I SEMESTER
BP805ET	PHARMACOVIGILANCE (THEORY)	45 Hours
of pharmac scenario of an organiz	s paper will provide an opportunity for the student to learn about covigilance as a science, basic terminologies used in pharmacovigi Pharmacovigilance, train students on establishing pharmacovigi ation, various methods that can be used to generate safety This paper also develops the skills of classifying drugs, diseases a	ilance, the global lance program in data and signal
	COURSE OUTCOMES	
CO 1	Understand pharmacovigilance fundamentals and ADR assessm	ent.
CO 2	Apply drug classification and coding for pharmacovigilance.	
CO 3	Implement and communicate pharmacovigilance methods.	
CO 4	Generate safety data and apply ICH guidelines.	
CO 5	Evaluate drug safety for diverse populations using pharmacogenomics.	
	COURSE CONTENT	
Unit I	Introduction to Pharmacovigilance	10 hours
	History and development of Pharmacovigilance	
	Importance of safety monitoring of Medicine	
	WHO International Drug Monitoring Programme	
	Pharmacovigilance Program of India (PvPI) Introduction to adverse design.	
	Introduction to adverse drug reactionsDefinitions and classification of ADRs	
	 Definitions and classification of ADRs Detection and reporting 	
	Methods in Causality assessment	
	Severity and seriousness assessment	
	Predictability and preventability assessment	
	Management of adverse drug reactions	
	Basic terminologies used in pharmacovigilance	
	Terminologies of adverse medication related events	
	Regulatory terminologies	
Unit II	Drug and disease classificationAnatomical, therapeutic and chemical classification of drugs	10

• International classification of diseases

• WHO adverse reaction terminologies

• International Non-proprietary Names for drugs Drug dictionaries and coding in pharmacovigilance

MedDRA and Standardized MedDRA queries

• Daily defined doses

WHO drug dictionary

hours

	,	
	Eudravigilance medicinal product dictionary	
	Information resources in pharmacovigilance	
	Basic drug information resources	
	Specialized resources for ADRs	
	Establishing pharmacovigilance program	
	Establishing in a hospital	
	Establishment & operation of drug safety department in the	
	industry	
	 Contract Research Organizations (CROs) 	
	Establishing a national programme	
Unit III	Vaccine safety surveillance	10 hours
	Vaccine Pharmacovigilance	20 110 011 0
	Vaccination failure	
	Adverse events following immunization	
	Pharmacovigilance methods	
	Passive surveillance – Spontaneous reports and case series	
	Stimulated reporting	
	• Active surveillance – Sentinel sites, drug event monitoring	
	and registries	
	 Comparative observational studies – Cross-sectional study, 	
	case-control study and cohort study	
	Targeted clinical investigations	
	Communication in pharmacovigilance	
	Effective communication in Pharmacovigilance	
	Communication in Drug Safety Crisis management	
	Communicating with Regulatory Agencies, Business	
	Partners, Healthcare facilities & Media	
Unit IV	Safety data generation	08 hours
	Pre-clinical phase	
	Clinical phase	
	 Post approval phase (PMS) 	
	ICH Guidelines for Pharmacovigilance	
	Organization and objectives of ICH	
	Expedited reporting	
	Individual case safety reports	
	Periodic safety update reports	
	Post approval expedited reporting	
	Pharmacovigilance planning	
	Good clinical practice in pharmacovigilance studies	
Unit V	Pharmacogenomics of adverse drug reactions	07 hours
	• Genetics related ADR with example focusing PK parameters.	
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Drug safety evaluation in special population

- Paediatrics
- Pregnancy and lactation
- Geriatrics

CIOMS

- CIOMS Working Groups
- CIOMS Form

CDSCO (India) and Pharmacovigilance

- D&C Act and Schedule Y
- Differences in Indian and global pharmacovigilance requirements

- 1. Textbook of Pharmacovigilance: S K Gupta, Jaypee Brothers, Medical Publishers.
- 2. Practical Drug Safety from A to Z By Barton Cobert, Pierre Biron, Jones and Bartlett Publishers.
- 3. Mann's Pharmacovigilance: Elizabeth B. Andrews, Nicholas, Wiley Publishers.
- 4. Stephens' Detection of New Adverse Drug Reactions: John Talbot, Patrick Walle, Wiley Publishers.
- 5. An Introduction to Pharmacovigilance: Patrick Waller, Wiley Publishers.
- 6. Cobert's Manual of Drug Safety and Pharmacovigilance: Barton Cobert, Jones& Bartlett Publishers.
- 7. Textbook of Pharmacoepidemiology edited by Brian L. Strom, Stephen E Kimmel, Sean Hennessy, Wiley Publishers.
- 8. A Textbook of Clinical Pharmacy Practice -Essential Concepts and Skills: G. Parthasarathi, Karin Northenden, Milap C. Nahata
- 9. National Formulary of India
- 10. Text Book of Medicine by Yashpal Munjal
- 11. Text book of Pharmacovigilance: concept and practice by GP Mohanta and PK Manna
- 12. http://www.whoumc.org/DynPage.aspx?id=105825&mn1=7347&mn2=7259&mn3=7297
- 13. http://www.ich.org/
- 14. http://www.cioms.ch/
- 15. http://cdsco.nic.in/
- 16. http://www.who.int/vaccine_safety/en/
- 17. http://www.ipc.gov.in/PvPI/pv_home.html

BP806ET	QUALITY CONTROL AND STANDARDIZATION OF HERBALS (THEORY)	45 Hours	
	Scope: In this subject the student learns about the various methods and guidelines fo		
	evaluation and standardization of herbs and herbal drugs. The subject also provides ar		
opportunit	y for the student to learn cGMP, GAP and GLP in traditional syster	n of medicines.	
	COURSE OUTCOMES		
CO 1	Learn the basics of WHO guidelines for making sure herbal drug high-quality.		
CO 2	Gain knowledge of the fundamental principles of quality assurant herbal drug industry.	ice (QA) in the	
CO 3	Develop the ability to comply with EU and ICH regulations gover quality, safety, and efficacy of herbal medicines.	ning the	
CO 4	Understand the principles and importance of stability testing to life and efficacy of herbal medicines.	ensure the shelf	
CO 5	Apply regulatory requirements and guidelines in the quality of products.	ontrol of herbal	
	COURSE CONTENT		
Unit I	Basic tests for drugs – Pharmaceutical substances, Medicinal	10 b	
	plants materials and dosage forms	10 hours	
	WHO guidelines for quality control of herbal drugs.		
	Evaluation of commercial crude drugs intended for use		
Unit II	Quality assurance in herbal drug industry of cGMP, GAP, GMP	10	
	and GLP in traditional system of medicine.		
	WHO Guidelines on Current Good Manufacturing Practices	hours	
	(cGMP) for Herbal Medicines		
	WHO Guidelines on GACP for Medicinal Plants.		
Unit III	EU and ICH guidelines for quality control of herbal drugs.	10 hours	
	Research Guidelines for Evaluating the Safety and Efficacy of Herbal Medicines		
Unit IV	Stability testing of herbal medicines. Application of various	08 hours	
	chromatographic techniques in standardization of herba		
	products.		
	Preparation of documents for new drug application and export	t	
	registration. GMP requirements and Drugs & Cosmetics Act		
	provisions.		
Unit V	Regulatory requirements for herbal medicines.	07 hours	
	WHO guidelines on safety monitoring of herbal medicines in	n	
	pharmacovigilance systems,		
	Comparison of various Herbal Pharmacopoeias.		
	Role of chemical and biological markers in standardization of	f	
	herbal products.		

- 1. Pharmacognosy by Trease and Evans
- 2. Pharmacognosy by Kokate, Purohit and Gokhale
- 3. Rangari, V.D., Text book of Pharmacognosy and Phytochemistry Vol. I, Carrier Pub., 2006.
- 4. Aggrawal, S.S., Herbal Drug Technology. Universities Press, 2002.
- 5. EMEA. Guidelines on Quality of Herbal Medicinal Products/Traditional Medicinal Products,
- 6. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation of Botanicals. Business Horizons Publishers, New Delhi, India, 2002.
- 7. Shinde M.V., Dhalwal K., Potdar K., Mahadik K. Application of quality control principles to herbal drugs. International Journal of Phytomedicine 1(2009); p. 4-8.
- 8. WHO. Quality Control Methods for Medicinal Plant Materials, World Health Organization, Geneva, 1998. WHO. Guidelines for the Appropriate Use of Herbal Medicines. WHO Regional Publications, Western Pacific Series No 3, WHO Regional office for the Western Pacific, Manila, 1998.
- 9. WHO. The International Pharmacopeia, Vol. 2: Quality Specifications, 3rd edn. World Health Organization, Geneva, 1981.
- 10. WHO. Quality Control Methods for Medicinal Plant Materials. World Health Organization, Geneva, 1999.
- 11. WHO. WHO Global Atlas of Traditional, Complementary and Alternative Medicine. 2 vol. set. Vol. 1 contains text and Vol. 2, maps. World Health Organization, Geneva, 2005.
- 12. WHO. Guidelines on Good Agricultural and Collection Practices (GACP) for Medicinal Plants. World Health Organization, Geneva, 2004.

BP807ET	COMPUTER AIDED DRUG DESIGN (THEORY)	45 Hours
_	s subject is designed to provide detailed knowledge of rational dru	g design process
and various techniques used in the rational drug design process.		
COURSE OUTCOMES		
CO1	Understand the principles and applications of computer-aid (CADD).	ed drug design
CO2	Analyze molecular modeling techniques for drug discovery and	optimization.
CO3	Evaluate structure-based and ligand-based drug design approac	hes.
CO4	Implement computational methods for predicting drug-target in	teractions.
CO5	Apply computer-aided drug design tools in rational drug discove	ery processes.
	COURSE CONTENT	
Unit I	Introduction to Drug Discovery and Development	101
	Stages of drug discovery and development	10 hours
	Lead discovery and Analog Based Drug Design	
	Rational approaches to lead discovery based on traditional	
	medicine, Random screening, Non-random screening,	
	serendipitous drug discovery, lead discovery based on drug	
	metabolism, lead discovery based on clinical observation.	
	Analog Based Drug Design: Bioisosterism, Classification, Bio	
	isosteric replacement. Any three case studies	
Unit II	Quantitative Structure-Activity Relationship (QSAR)	10
	SAR versus QSAR, History and development of QSAR, Types of	1
	physicochemical parameters, experimental and theoretical	
	approaches for the determination of physicochemical parameters such as Partition coefficient, Hammet's substituent	
	constant and Tafts steric constant. Hansch analysis, Free Wilson	
	analysis, 3D-QSAR approaches like COMFA and COMSIA.	
Unit III	Molecular Modeling and virtual screening techniques	10 hours
	Virtual Screening techniques: Drug likeness screening,	
	Concept of pharmacophore mapping and pharmacophore-based	
	Screening,	
	Molecular docking: Rigid docking, flexible docking, manual	
	docking, Docking based screening. De novo drug design.	
Unit IV	Informatics & Methods in Drug Design	08 hours
	Introduction to Bioinformatics, cheminformatics. ADME	
	databases, chemical, biochemical and pharmaceutical databases	
Unit V	Molecular Modeling: Introduction to molecular mechanics and	
	quantum mechanics. Energy Minimization methods and	
	Conformational Analysis, global conformational minima	l

determination.

- 1. Robert GCK, ed., "Drug Action at the Molecular Level" University Prak Press Baltimore.
- 2. Martin YC. "Quantitative Drug Design" Dekker, New York.
- 3. Delgado JN, Remers WA eds "Wilson & Gisvolds's Text Book of Organic Medicinal & Pharmaceutical Chemistry" Lippincott, New York.
- 4. Foye WO "Principles of Medicinal chemistry 'Lea & Febiger.
- 5. Koro lkovas A, Burckhalter JH. "Essentials of Medicinal Chemistry" Wiley Interscience.
- 6. Wolf ME, ed "The Basis of Medicinal Chemistry, Burger's Medicinal Chemistry" John Wiley & Sons, New York.
- 7. Patrick Graham, L., An Introduction to Medicinal Chemistry, Oxford University Press.
- 8. Smith HJ, Williams H, eds, "Introduction to the principles of Drug Design" Wright Boston.
- 9. Silverman R.B. "The organic Chemistry of Drug Design and Drug Action" Academic Press New York.

BP808ET	CELL AND MOLECULAR BIOLOGY (THEORY)	45 Hours
Scope:		

- Cell biology is a branch of biology that studies cells their physiological properties, their structure, the organelles they contain, interactions with their environment, their life cycle, division, death and cell function.
- This is done both on a microscopic and molecular level.
- Cell biology research encompasses both the great diversity of single-celled organisms like bacteria and protozoa, as well as the many specialized cells in multi-cellular organisms such as humans, plants, and sponges.

such as numans, plants, and sponges.		
COURSE OUTCOMES		
CO1	Understand the fundamental concepts and history of cell and molecular biology,	
	including cell properties and reproduction.	
CO2	Comprehend DNA and RNA roles, types, and the processes of tra- translation.	•
СО3	Understand protein structure, synthesis, and their role in cellular pathways.	processes and
CO4	Analyze genetic principles, the cell cycle, and the processes meiosis.	of mitosis and
CO5	Understand cell signaling mechanisms, receptor functions, and the impact of signaling pathway misregulation.	
COURSE CONTENT		
Unit I	a) Cell and Molecular Biology: Definitions theory and basics and	10 hours
	Applications.	10 110013
	b) Cell and Molecular Biology: History and Summation.	
	c) Properties of cells and cell membrane.	
	d) Prokaryotic versus Eukaryotic	
	e) Cellular Reproduction	
	f) Chemical Foundations – an Introduction and Reactions (Types)	
Unit II	a) DNA and the Flow of Molecular Information	10
	b) DNA Functioning	1
	c) DNA and RNA	hours
	d) Types of RNA	
	e) Transcription and Translation	
Unit III	a) Proteins: Defined and Amino Acids	10 hours
	b) Protein Structure	
	c) Regularities in Protein Pathways	
	d) Cellular Processes	
	e) Positive Control and significance of Protein Synthesis	
Unit IV	a) Science of Genetics	08 hours
	b) Transgenics and Genomic Analysis	

	c) Cell Cycle analysis	
	d) Mitosis and Meiosis	
	e) Cellular Activities and Checkpoints	
Unit V	a) Cell Signals: Introduction	07 hours
	b) Receptors for Cell Signals	
	c) Signaling Pathways: Overview	
	d) Misregulation of Signaling Pathways	
	e) Protein-Kinases: Functioning	

- 1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.
- 2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.
- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. Edward: Fundamentals of Microbiology.
- 10. N.K.Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 11. Bergey manual of systematic bacteriology, Williams and Wilkins- A Waverly company
- 12. B.R. Glick and J.J. Pasternak: Molecular Biotechnology: Principles and Applications of recombinant DNA: ASM Press Washington D.C.
- 13. RA Goldshy et. al., : Kuby Immunology.

BP809ET	COSMETIC SCIENCE (THEORY)	45 Hours	
Scope:	Scope:		
	COURSE OUTCOMES		
CO1	Classify cosmetic products based on regulatory definitions, explain the evolution of cosmeceuticals, and categorize cosmetic excipients by their functions.		
COI			
CO2	Analyze the formulation principles and advantages/disadvantage	ges of skin care,	
COZ	hair care, and oral care products.		
	Evaluate sunscreens based on SPF classification, assess the r		
CO3	cosmetic formulations, and apply BIS specifications and analyti	cal methods for	
	cosmetic products.		
CO4	Assess cosmetic efficacy using tools like sebumeter and corneo	meter, measure	
	parameters and evaluate the benefits of soaps and syndet bars.	l aggagg alsin agna	
CO5	Analyze causes of oily and dry skin, evaluate cosmetic issues and	assess skincare	
	concerns.		
	COURSE CONTENT		
Unit I	Classification of cosmetic and cosmeceutical products	_	
Onici	Definition of cosmetics as per Indian and EU regulations,	10 hours	
	Evolution of cosmeceuticals from cosmetics, cosmetics as quasi		
	and OTC drugs		
	Cosmetic excipients: Surfactants, rheology modifiers,		
	humectants, emollients, preservatives. Classification and		
	application		
	Skin : Basic structure and function of skin.		
	Hair: Basic structure of hair. Hair growth cycle.		
	Oral Cavity : A common problem associated with teeth and gums.		
Unit II	Principles of formulation and building blocks of skin care	10	
	products:	hours	
	Face wash, Moisturizing cream, Cold Cream, Vanishing cream and their advantages and disadvantages. Application of these		
	products in the formulation of cosmeceuticals.		
	Antiperspirants & deodorants - Actives & mechanism of action.		
	Principles of formulation and building blocks of Hair care		
	products: Conditioning shampoo, Hair conditioner, anti-		
	dandruff shampoo, Hair oils.		
	Chemistry and formulation of Para-phenylene diamine-based		
	hair dye.		
	Principles of formulation and building blocks of oral care		
	products:		
	Toothpaste for bleeding gums, and sensitive teeth. Teeth		
	whitening, Mouthwash.		

Unit III	Sun protection, Classification of Sunscreens and SPF.	10 hours
	Role of herbs in cosmetics:	10 110415
	Skin Care: Aloe and turmeric	
	Hair care: Henna and amla.	
	Oral care: Neem and clove	
	Analytical cosmetics: BIS specification and analytical methods	
	for shampoo, skin cream and toothpaste.	
Unit IV	Principles of Cosmetic Evaluation: Principles of sebumeter,	08 hours
	corneometer. Measurement of TEWL, Skin Color, Hair tensile	
	strength, Hair combing properties Soaps, and syndet bars.	
	Evolution and skin benefits.	
Unit V	Oily and dry skin, causes leading to dry skin, skin moisturization.	07 hours
	Basic understanding of the terms Comedogenic, and dermatitis.	
	Cosmetic problems associated with Hair and scalp: Dandruff,	
	Hair fall causes	
	Cosmetic problems associated with skin: blemishes, wrinkles,	
	acne, prickly heat and body odor.	
	Antiperspirants and Deodorants- Actives and mechanism of	
	action	

- 1. Harry's Cosmeticology, Wilkinson, Moore, Seventh Edition, George Godwin.
- 2. Cosmetics Formulations, Manufacturing and Quality Control, P.P. Sharma, 4th Edition, Vandana Publications Pvt. Ltd., Delhi.
- 3. Text book of cosmeticology by Sanju Nanda & Roop K. Khar, Tata Publishers.

	DIVADIVAÇÕI OCICAL CODERNING MEMILODO	
BP810ET	PHARMACOLOGICAL SCREENING METHODS (THEORY)	45 Hours
Scope: Thi	is subject is designed to impart the basic knowledge of precli	nical studies in
experimental animals including design, conduct and interpretations of results.		
	COURSE OUTCOMES	
CO1	CO1 Understand guidelines and techniques for using laboratory animals.	
CO2	Apply models to evaluate CNS drug effects.	
CO3	Use models to study ANS drug effects.	
CO4	Evaluate models for CVS and other drug effects.	
CO5	Develop research skills and analyze data statistically.	
	COURSE CONTENT	
Unit I	Laboratory Animals:	08 hours
	Study of CPCSEA and OECD guidelines for maintenance, breeding	
	and conduct of experiments on laboratory animals, Common lab	
	animals: Description and applications of different species and	
	strains of animals. Popular transgenic and mutant animals.	
	Techniques for collection of blood and common routes of drug	
	administration in laboratory animals, Techniques of blood	
Unit II	collection and euthanasia.	
Unitn	Preclinical screening modelsa. Introduction: Dose selection, calculation and conversions,	10 hours
	preparation of drug solution/suspensions, grouping of animals	
	and importance of sham negative and positive control groups.	
	Rationale for selection of animal species and sex for the study.	
	b. Study of screening animal models for Diuretics, nootropics,	
	anti-Parkinson's, antiasthmatics,	
	Preclinical screening models: for CNS activity- analgesic,	
	antipyretic, anti-inflammatory, general anesthetics, sedative and	
	hypnotics, antipsychotics, antidepressants, antiepileptics, anti-	
	parkinsonism, Alzheimer's disease	
Unit III	Preclinical screening models: for ANS activity,	10 Hours
	sympathomimetics, sympatholytic, parasympathomimetics, parasympatholytic, skeletal muscle relaxants, drugs acting on	
	eye, local anesthetics	
Unit IV	Preclinical screening models: for CVS activity	10 hours
	antihypertensives, diuretics, antiarrhythmic, antidyslepidemic	,
	anti-aggregatory, coagulants, and anticoagulants	
	Preclinical screening models for other important drugs like	
	antiulcer, antidiabetic, anticancer and antiasthmatics.	0=1
Unit V	Research methodology and Bio-statistics	07 hours

Selection of research topic, review of literature, research	
hypothesis and study design	
Pre-clinical data analysis and interpretation using Students 't'	
test and One-way ANOVA. Graphical representation of data	

- 1. Fundamentals of experimental Pharmacology-byM.N.Ghosh
- 2. Hand book of Experimental Pharmacology-S.K.Kulkarni
- 3. CPCSEA guidelines for laboratory animal facility.
- 4. Drug discovery and Evaluation by Vogel H.G.
- 5. Drug Screening Methods by Suresh Kumar Gupta and S. K. Gupta
- 6. Introduction to biostatistics and research methods by PSS Sundar Rao and J Richard

DD044EF	ADVANCED INSTRUMENTATION TECHNIQUES	45 Hours
BP811ET	(THEORY)	10 110410

Scope: This subject deals with the application of instrumental methods in qualitative and quantitative analysis of drugs. This subject is designed to impart advanced knowledge on the principles and instrumentation of spectroscopic and chromatographic hyphenated techniques. This also emphasizes on theoretical and practical knowledge on modern

analytical instruments that are used for drug testing.		
	COURSE OUTCOMES	
CO1	Understand principles, instrumentation, and applications of NMR and Mass Spectrometry.	
CO2	Analyze thermal and X-ray diffraction methods, including their principles, instrumentation, and applications.	
CO3	Perform calibration and validation of various analytical instrument and USFDA guidelines.	ents as per ICH
CO4	Apply principles and techniques of radioimmune assays and extraction methods for drug analysis.	
CO5	Utilize advanced hyphenated techniques (LC-MS/MS, GC-MS/MS,	HPTLC-MS) for
COS	qualitative and quantitative drug analysis.	
COURSE CONTENT		
Unit I	Nuclear Magnetic Resonance spectroscopy	10 hours
	Principles of H-NMR and C-NMR, chemical shift, factors affecting	
	chemical shift, coupling constant, Spin - spin coupling, relaxation,	
	instrumentation and applications	
	Mass Spectrometry- Principles, Fragmentation, Ionization	
	techniques –Electron impact, chemical ionization, MALDI, FAB,	
	Analyzers-Time of flight and Quadrupole, instrumentation,	
	applications	
Unit II	Thermal Methods of Analysis : Principles, instrumentation and	10 hours
	applications of thermogravimetric analysis (TGA), Differential	
	Thermal Analysis (DTA), Differential Scanning Calorimetry	
	(DSC)	
	X-Ray Diffraction Methods: Origin of X-rays, basic aspects of	
	crystals, Xray Crystallography, rotating crystal technique, single	
	crystal diffraction, powder diffraction, structural elucidation and	
	applications.	
Unit III	Calibration and validation-as per ICH and USFDA guidelines	10 hours
	Calibration of following Instruments Electronic balance, UV-Visible spectrophotometer, IR	
	Lieutonic balance, ov-visible spectrophotometer, ik	

spectrophotometer Fluorimeter, Flame Photometer, HPLC and

Importance,

GC

Radio

immunoassay:

Unit IV

08 hours

components,

various

	Principle, different methods, Limitation and Applications of	
	Radio immunoassay	
	Extraction techniques: General principle and procedure	
	involved in the solid phase extraction and liquid-liquid extraction	
Unit V	Hyphenated techniques -LC-MS/MS, GC-MS/MS, HPTLC-MS.	07 hours

- 1. Instrumental Methods of Chemical Analysis by B.K Sharma
- 2. Organic spectroscopy by Y.R Sharma
- 3. Text book of Pharmaceutical Analysis by Kenneth A. Connors
- 4. Vogel's Text book of Quantitative Chemical Analysis by A.I. Vogel
- 5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake
- 6. Organic Chemistry by I. L. Finar
- 7. Organic spectroscopy by William Kemp
- 8. Quantitative Analysis of Drugs by D. C. Garrett
- 9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi
- 10. Spectrophotometric identification of Organic Compounds by Silverstein

BP812ET	DIETARY SUPPLEMENTS AND NUTRACEUTICALS (THEORY)	45 Hours
_	s subject covers foundational topic that are important for unders	-
and require	ements of dietary supplements among different groups in the pop	ulation.
	COURSE OUTCOMES	
CO1	Develop a clear understanding of functional foods, nutraceutical supplements, including their definitions and distinctions.	
Gain a comprehensive understanding of phytochemicals as bioactive Compounds found in plants, distinct from traditional nutrients.		
Learn a comprehensive understanding of free radicals and ROS as with unpaired electrons, leading to high reactivity.		
Gain a comprehensive understanding of free radicals, their form effects, and implications across various health and disease conte		exts.
CO5	Apply regulatory knowledge to assess food safety practi processing, and distribution.	
COURSE CONTENT		
Unit I	a. Definitions of Functional foods, Nutraceuticals and Dietary supplements. Classification of Nutraceuticals, Health problems and diseases that can be prevented or cured by Nutraceuticals i.e. weight control, diabetes, cancer, heart disease, stress, osteoarthritis, hypertension etc. b. Public health nutrition, maternal and child nutrition, nutrition and ageing, nutrition education in the community. c. Source, Name of marker compounds and their chemical nature, Medicinal uses and health benefits of the following used as nutraceuticals/functional foods: Spirulina, Soybean, Ginseng, Garlic, Broccoli, Gingko, Flaxseeds	o nours
Unit II	 Phytochemicals as nutraceuticals: Occurrence and characteristic features (chemical nature medicinal benefits) of following a) Carotenoids- α and β-Carotene, Lycopene, Xanthophylls, leuting b) Sulfides: Diallyl sulfides, Allyl trisulfide. c) Polyphenolics: Reservetrol d) Flavonoids- Rutin, Naringin, Quercitin, Anthocyanidins, catechins, Flavones e) Prebiotics/Probiotics.: Fructooligosaccharides, Lactobacillus f) Phyto estrogens: Isoflavones, daidzein, Geebustin, lignans g) Tocopherols h) Proteins, vitamins, minerals, cereal, vegetables and beverages as functional foods: oats, wheat bran, rice bran, sea foods, coffee, tea and the like. 	15 nours

Unit III	 a) Introduction to free radicals: Free radicals, reactive oxygen species, production of free radicals in cells, damaging reactions of free radicals on lipids, proteins, Carbohydrates, and nucleic acids. b) Dietary fibres and complex carbohydrates as functional food ingredients. 	07 hours
Unit IV	 a) Free radicals in Diabetes mellitus, Inflammation, Ischemic reperfusion injury, Cancer, Atherosclerosis, Free radicals in brain metabolism and pathology, kidney damage, muscle damage. Free radicals' involvement in other disorders. Free radicals' theory of ageing. b) Antioxidants: Endogenous antioxidants – enzymatic and nonenzymatic antioxidant defence, Superoxide dismutase, catalase, Glutathione peroxidase, Glutathion Vitamin C, Vitamin E, α- Lipoic acid, melatonin Synthetic antioxidants: Butylated hydroxy Toluene, Butylated hydroxy Anisole. c) Functional foods for chronic disease prevention 	10 hours
Unit V	 a) Effect of processing, storage and interactions of various environmental factors on the potential of nutraceuticals. b) Regulatory Aspects; FSSAI, FDA, FPO, MPO, AGMARK. HACCP and GMPs on Food Safety. Adulteration of foods. c) Pharmacopoeial Specifications for dietary supplements and nutraceuticals. 	06 hours

- 1. Dietetics by Sri Lakshmi
- 2. Role of dietary fibres and neutraceuticals in preventing diseases by K. T Agusti and P. Faizal: BSPunblication.
- 3. Advanced Nutritional Therapies by Cooper. K.A., (1996).
- 4. The Food Pharmacy by Jean Carper, Simon & Schuster, UK Ltd., (1988).
- 5. Prescription for Nutritional Healing by James F. Balch and Phyllis A. Balch 2nd Edn., Avery Publishing Group, NY (1997).
- 6. G. Gibson and C. Williams Editors 2000 Functional foods Woodhead Publ.Co. London.
- 7. Goldberg, I. Functional Foods. 1994. Chapman and Hall, New York.
- 8. Labuza, T.P. 2000 Functional Foods and Dietary Supplements: Safety, Good Manufacturing Practice (GMPs) and Shelf-Life Testing in Essentials of Functional Foods M.K. Sachmidl and T.P. Labuza eds. Aspen Press.
- 9. Handbook of Nutraceuticals and Functional Foods, Third Edition (Modern Nutrition)
- 10. Shils, ME, Olson, JA, Shike, Shike, M. 1994 Modern Nutrition in Health and Disease. Eighth edition. Lea and Febiger

BP813ET	Pharmaceutical Product Development (THEORY)	45 Hours		
-	s subject covers foundational topic that are important for unders	-		
and require	ements of dietary supplements among different groups in the pop	ulation.		
COURSE OUTCOMES				
	Outline the objectives and regulations governing pre-formulat			
CO1	development, stability assessment, manufacturing, and quality of	control testing of		
	various dosage forms. Learners will demonstrate specialized knowledge of pharmaceu			
CO2				
Analyze the tablet, capsule, parenteral, aerosol products and novel drug systems (NDDS) excipients.		vel drug delivery		
CO4	Apply various optimization techniques such as factorial design	• • •		
601	Design (QbD) principles in pharmaceutical product developmen			
CO5	Assess the selection and quality control testing of packagin	Ö		
	pharmaceutical products, considering regulatory considerations	i.		
	COURSE CONTENT			
Unit I	Introduction to pharmaceutical product development,	107		
011101	objectives, regulations related to preformulation, formulation	10 Hours		
	development, stability assessment, manufacturing and quality			
	control testing of different types of dosage forms			
Unit II	An advanced study of Pharmaceutical Excipients in	10 hours		
	pharmaceutical product development	10 Hours		
	with a special reference to the following categories			
	i. Solvents and solubilizers			
	ii. Cyclodextrins and their applications			
	iii. Non-ionic surfactants and their applications			
	iv. Polyethylene glycols and sorbitol			
	v. Suspending and emulsifying agent			
Unit III	vi. Semi solid excipients An advanced study of Pharmacoutical Excipients in			
UIIIL III	An advanced study of Pharmaceutical Excipients in pharmaceutical product development with a special reference to	10 Hours		
	the following categories			
	i. Tablet and capsule excipients			
	ii. Directly compressible vehicles			
	iii. Coat materials			
	iv. Excipients in parenteral and aerosols products			
	v. Excipients for formulation of NDDS			
	Selection and application of excipients in pharmaceutical			
	formulations with specific industrial applications			
Unit IV	Optimization techniques in pharmaceutical product			
	development. A study of various optimization techniques for	•		

	pharmaceutical product development with specific examples.	
	Optimization by factorial designs and their applications. A study	
	of QbD and its application in pharmaceutical product	
	development.	
Unit V	Selection and quality control testing of packaging materials for	07 hours
	pharmaceutical product development- regulatory	
	considerations.	

- 1. Pharmaceutical statistics practical and clinical applications by Stanford Bolton, Charlesbon; Marcel Dekker inc.
- 2. Encyclopedia of pharmaceutical technology, edited by James Swarbrick, third Edition, Informa healthcare publishers.
- 3. Pharmaceutical dosage forms, tablets, volume ii, edited by Herbert a. Lieberman Andleon Lachman; Marcel Dekker, inc.
- 4. The theory and practice of industrial pharmacy, fourth edition, edited by Roop Khar, S P Vyas, Farhan J Ahmad, Gaurav K Jain; CBS publishers and distributors Pvt. ltd. 2013.
- 5. Martin's physical pharmacy and pharmaceutical sciences, fifth edition, edited by Patrick j. Sinko, bi publications pvt. Ltd.
- 6. Targeted and controlled drug delivery, novel carrier systems by s. P. Vyas and R. K. Khar, CBS publishers and distributors pvt. Ltd, first edition 2012.
- 7. Pharmaceutical dosage forms and drug delivery systems, Loyd v. Allen jr., Nicholas B. POPOVICH, Howard c. Ansel, 9th ed. 40
- 8. Aulton's pharmaceutics the design and manufacture of medicines, Michael e. Aulton, 3rd ed.
- 9. Remington the science and practice of pharmacy, 20th ed.
- 10. Pharmaceutical dosage forms tablets vol 1 to 3, a. Liberman, Leon Lachman Andjoseph B. Schwartz
- 11. Pharmaceutical dosage forms disperse systems vol 1 to 3, H.A. Liberman, Martin, M.R and Gilbert S. Banker.
- 12. Pharmaceutical dosage forms parenteral medication vol 1 & 2, Kenneth e. Avis Andh.A. Libermann.
- 13. Advanced review articles related to the topics